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- (71) Applicant (*for all designated States except US*): **GLAXO GROUP LIMITED** [GB/GB]; Glaxo Wellcome House, Berkeley Avenue, Greenford, Middlesex UB6 ONN (GB).
- (72) Inventors; and
- (75) Inventors/Applicants (*for US only*): **DAVIS, Stephen, Thomas** [US/US]; GlaxoSmithKline, Five Moore Drive, P.O. Box 13398, Research Triangle Park, NC 27709 (US). **DEV, Inderjit, Kumar** [US/US]; GlaxoSmithKline, Five Moore Drive, P.O. Box 13398, Research Triangle Park, NC 27709 (US). **DORNSIFE, Ronna, Ellen** [US/US]; GlaxoSmithKline, Five Moore Drive, P.O. Box 13398, Research Triangle Park, NC 27709 (US). **GRAY, John, Gordon** [US/US]; GlaxoSmithKline, Five Moore Drive, P.O. Box 13398, Research Triangle Park, NC 27709 (US).
- (74) Agents: **LEMANOWICZ, John, L.** et al.; GlaxoSmithKline, Five Moore Drive, P.O. Box 13398, Research Triangle Park, NC 27709 (US).
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(54) Title: CANCER TREATMENT METHOD

(57) Abstract: The invention relates to a method for treating cancer including administering to a mammal therapeutically effective amounts of an anti-neoplastic agent and a PDE4 inhibitor, treatment combinations containing the same as well as a method for attenuating unwanted cancer treatment side effects including myelosuppression and mucositis.



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CANCER TREATMENT METHOD

BACKGROUND OF THE INVENTION

5 The present invention relates to a method of treating cancer in a mammal and to pharmaceutical combinations useful in such treatment. In particular, the method relates to a cancer treatment method that includes administering anti-neoplastic agents which adversely effect normal progenitor cells and phosphodiesterase 4 (PDE4) inhibitors to the mammal. Such PDE4 inhibitors attenuate unwanted side effects on
10 normal progenitor cells of the mammal resulting from administration of said anti-neoplastic agent.

 Effective chemotherapy for cancer treatment, which has acceptable toxicity to normal cells, is a continuing goal in the oncology field. Numerous agents are used in
15 the treatment of cancer, including agents that adversely affect normal progenitor cells upon use. One such class of anti-neoplastic agents that have an adverse effect on normal progenitor cells is known as phase specific anti-neoplastic agents. Phase specific anti-neoplastic agents have activity against cells that are in the process of cell division. Typically, such agents have effect on the cell cycle at G₁ – the period
20 between mitosis and DNA synthesis; S – the period of DNA synthesis; G₂ – the pre-mitotic interval; and/or M – the period of mitosis. These agents are not effective in G₀, the quiescent or resting cell phase. Therefore, phase specific anti-neoplastic agents are active against cells in the process of cell division. Phase specific anti-neoplastic agents are most effective against cancers that have a large growth fraction,
25 that is, tumors that have a high percentage of dividing cells. Problematically, however, such agents also have an adverse effect on rapidly proliferating, normal tissues such as bone marrow, hair follicles and intestinal epithelium. (See Goodman & Gilman's, *The Pharmacologic Basis Of Therapeutics* 9th Ed., pages 1230-1232.)

30 As indicated above, the bone marrow and epithelial mucosa (gastrointestinal and oral) have a rapid cell cycle and are particularly sensitive to phase specific anti-

neoplastic agents. Consequently, myelosuppression and mucositis are the two most common dose-limiting side effects of anti-neoplastic treatment. Myelosuppression is suppression of normal bone marrow activity (hematopoiesis) that ultimately causes a reduction in white blood cells (neutropenia), platelets (thrombocytopenia), and/or red blood cells (anemia) in the peripheral blood. All of these adverse effects can lead to complications to the patient's health. For instance, neutropenia can lead to susceptibility to recurrent and life threatening, opportunistic, bacterial infections. Mucositis is inflammation resulting from damage to the epithelial mucosae. Mucositis induced by anti-neoplastic therapies includes damage to the epithelial barrier and potential entry of microbial flora into the blood. Neutropenia and mucositis, if severe enough, limit patients' ability to tolerate subsequent courses of anti-neoplastic drugs (Kim SK, Demitri GD, Hematology/Oncology Clinics of North America, 10 (2) 377-393, (1996)).

While effective treatment for mucositis is still a major unmet medical need and continues to be a threat to patients treated with anti-neoplastic agents, several techniques have been utilized to reduce and/or control myelosuppression. Patients with neutropenia are commonly treated with rescue agents such as granulocyte-colony stimulating factor (G-CSF) to reduce the duration and severity of neutropenia. Although this approach has been proven to reduce peripheral neutropenia, no survival benefit to patients from either reduced infection or dose-intensification of chemotherapy has been demonstrated (Higa, GD; Pharmacotherapy, 18 (6) 1249-1254, (1998); Johnston, E; Crawford, J; Seminars in Oncology, 25 (5): 552-561, (1998); Petros, WP; Crawford, J; Current Opinion in Hematology, 4 (3): 213-216). Moreover, recent evidence suggests that stimulation by G-CSF forces surviving stem cells into proliferation and repeated rounds of chemotherapy further damages and depletes stem cell pools. (van Os, R et al., Blood, 92 (6), 1950-1956, (1997); Higa, GD; Pharmacotherapy, 18 (6) 1249-1254, (1998); Johnston, E; Crawford, J; Seminars in Oncology, 25 (5) 552-561, (1998)).

Instead of stimulating proliferation of damaged and/or surviving stem cells, a strategy of total or partial preservation of normal cells by chemoprotection would be a superior approach. Many investigators have suggested that stem cells can be protected from radiation and chemotherapy damage by keeping the stem cells in a quiescent state during drug exposure. Numerous studies have demonstrated that elevated concentration of intracellular cAMP in certain cell types result in suppression of cell proliferation and activation. (Vairo, G et al., J Biol Chem, 265 (6): 2692-2700, (1990); Berridge, MV et al., Experimental Hematology, 21: 269-276 (1993); Morley, A et al., Proc Soc Exp Biol Med, 138: 57-59, (1997); Ralph, RK, FEBS Lett, 161: 1-8). Increased intracellular cAMP also increases the survival and half-life of mature peripheral blood neutrophils apparently by decreasing natural apoptosis (Walker, BA et al., J Immunol, 158: 2926-2931; Rossi, AG, Biochem Biophys Research Commun, 217 (3), 892-899 (1995); Niwa, M et al. Euro J Pharmacol 371, 59-67, (1999)).

The present inventors hypothesized that elevated levels of intracellular cAMP might protect normal progenitor and stem cells in various tissues from the cytotoxicities produced by anti-neoplastic agents through induction of cell cycle arrest and/or by inhibition of apoptosis. Intracellular concentrations of cAMP can be elevated by several mechanisms (see Figure 1), including activating surface receptors which are G-protein coupled to the cAMP generating enzyme adenylyl cyclase (e.g. prostaglandin E-2 [PGE₂]) 1; increasing cAMP levels by directly stimulating adenylyl cyclase (e.g. forskolin) 2; by treating with a stable analog of cAMP (e.g., dibutyl cAMP, 8-Cl cAMP, or 8-bromo cAMP) 3, or by inhibiting cyclic nucleotide phosphodiesterases (PDEs), which decreases the rate at which cAMP is catabolized to AMP 4.

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Phosphodiesterases (PDEs) comprise a large, divergent family of enzymes that catalyze the catabolism of cAMP and cGMP to AMP and GMP, respectively. Eleven members of this family have been identified to date based on substrate specificity, kinetic properties, sensitivity to specific inhibitors, tissue distribution, and primary sequences (Manganiello, VC et al., Arch Biochem Biophys, 322 (1): 1-13, (1995);

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Fawcett, L et al., Proc Natl Acad Sci USA, 97(7): 3702-3707 (2000)). Recognition of these differences has greatly stimulated interest in PDEs as drug targets. In particular, the cAMP specific PDE, termed PDE4, which is found in nearly all immune and inflammatory cells is an attractive target for novel anti-asthmatic and anti-inflammatory therapies.

The present inventors have now discovered a new method of treating cancer using a novel pharmaceutical combination, which can selectively treat cancers, while attenuating the associated adverse, side effects including myelosuppression and mucositis. The present invention recognizes the use of PDE4 inhibitors as selective agents that could provide protection from myelosuppression and mucositis induced by anti-neoplastic agents without affecting the anti-neoplastic efficacy of these drugs. Elevation of cAMP by inhibiting PDE4 isoenzyme offers a definite advantage over inhibiting multiple PDE isoenzymes because it can potentially lead to an increase of intracellular cAMP in normal cells, but not in tumor cells. PDE4 is the exclusive cAMP PDE isoenzyme found in neutrophils and is the predominant cAMP PDE isoenzyme in epithelial cells. Selective inhibition of PDE4 in neutrophils or epithelial cells would raise the intracellular level of cAMP in those cells significantly. In contrast, tumor cells have other cAMP PDE isoenzymes in addition to PDE4. These additional isoenzymes in tumor cells would allow cAMP to be catabolized to AMP and thus prevent a buildup of intracellular cAMP even if treated with a selective PDE4 inhibitor.

SUMMARY OF THE INVENTION

In a first aspect of the present invention, there is provided a method of treating cancer in a mammal, including: administering to said mammal therapeutically effective amounts of (i) an anti-neoplastic agent which has an adverse effect on normal progenitor cells upon use; and (ii) a PDE4 inhibitor.

In a second aspect of the present invention, there is provided a method of treating cancer in a mammal, including: administering to said mammal therapeutically

effective amounts of (i) paclitaxel; and (ii) *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.

5 In a third aspect of the present invention, there is provided a method of treating cancer in a mammal, including: administering to said mammal therapeutically effective amounts of (i) topotecan; and (ii) *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.

10 In a fourth aspect of the present invention, there is provided a cancer treatment combination, including: therapeutically effective amounts of (i) an anti-neoplastic agent which has an adverse effect on normal progenitor cells upon use; and (ii) a PDE4 inhibitor.

15 In a fifth aspect of the present invention, there is provided a cancer treatment combination, including: therapeutically effective amounts of (i) paclitaxel; and (ii) *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.

20 In a sixth aspect of the present invention, there is provided a cancer treatment combination, including: therapeutically effective amounts of (i) topotecan; and (ii) *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.

25 In a seventh aspect of the present invention, there is provided a method of attenuating, in a mammal, cancer treatment side effects on normal progenitor cells in the mammal, including: administering to the mammal a therapeutically effective amount of a PDE4 inhibitor.

30 In an eighth aspect of the present invention, there is provided use of (i) an anti-neoplastic agent which has an adverse effect on normal progenitor cells upon use; and (ii) a PDE4 inhibitor in therapy.

In a ninth aspect of the present invention, there is provided use of (i) an anti-neoplastic agent which has an adverse effect on normal progenitor cells upon use; and (ii) a PDE4 inhibitor in the preparation of a medicament for the treatment of cancer.

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In a tenth aspect of the present invention, there is provided use of a PDE4 inhibitor in the preparation of a medicament for the attenuation of cancer treatment side effects on normal progenitor cells.

10 In an eleventh aspect of the present invention, there is provided a method of treating cancer in a mammal, including administering to the mammal a therapeutically effective amount of a PDE4 inhibitor.

BRIEF DESCRIPTION OF THE DRAWINGS

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Figure 1 depicts various mechanisms for increasing intracellular concentrations of cAMP in cells.

20 Figure 2 depicts a flow cytometric evaluation of expanded CD34⁺ stem cells treated with (a) DMSO; (b) rolipram plus forskolin followed by treatment with various doses of paclitaxel; and (c) a plot of percent live cells vs. paclitaxel dose with pre-treatment with DMSO or rolipram plus forskolin.

25 Figure 3 (a), (b), and (c) depict protection of specific-stage neutrophil progenitors (CFU-GMs) as measured by colony forming assay wherein cells are treated with rolipram plus forskolin followed by treatment with either (a) 5-fluorouracil, (b) etoposide, or (c) paclitaxel.

30 Figure 4 (a), (b), (c), (d), (e), (f), and (g) depict protection of multiple-stage neutrophil progenitors as measured by a CD15 neutrophil antigen-cell-based ELISA (CELISA) method wherein cells are treated with rolipram plus forskolin followed by

treatment with either (a) paclitaxel, (b) 5-fluorouracil, (c) etoposide, (d) vinblastine, (e) vinorelbine, (f) methotrexate, or (g) topotecan.

Figure 5 (a), (b), (c), (d), (e), (f), (g), (h), and (i) depict protection of multiple-stage neutrophil progenitors as measured by a CD15 neutrophil antigen-cell-based ELISA (CELISA) method wherein cells are treated with (a) Ariflo, (b) Piclamilast, (c) CDP-840 [GR259653X], (d) CDP-840 [GR259654X], (e) Nitraquazone, (f) Denbufylline, (g) RS-25344, (h) CP-77059 or (i) GI193600X, each plus forskolin followed by treatment with paclitaxel.

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Figure 6 depicts protection of multiple-stage neutrophil progenitors as measured by a CD15 neutrophil antigen-cell-based ELISA (CELISA) method wherein cells are treated with ariflo plus forskolin followed by treatment with topotecan.

Figure 7 (a), (b), and (c) depict non-protection of multiple-stage neutrophil progenitors as measured by a CD15 neutrophil-cell-based ELISA (CELISA) method wherein cells are treated with rolipram plus forskolin followed by treatment with either (a) cisplatin, (b) doxorubicin, or (c) carboplatin.

Figure 8 (a), (b), (c), (d), (e), (f), (g), and (h) depict no-effect on the anti-neoplastic effect of paclitaxel against either the human RKO colon tumor line ((a) and (c)) or the human HT-29 colon tumor line ((b) and (d)), or the human MDA468 breast tumor line ((e) and (g)) or the human A549 lung tumor line ((f) and (h)) wherein relative growth of tumor cells is measured with the protein stain sulforhodamine (SRB) after cells are treated with rolipram plus forskolin or Ariflo plus forskolin followed by treatment with 100 nM paclitaxel.

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Figure 9 (a) and (b) depict PDE4 inhibitor compounds, (a) ariflo and (b) piclamilast, useful in the present invention.

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Figure 10 (a) and (b) depict PDE4 inhibitor compounds, (a) filaminast and (b) CDP-840, useful in the present invention.

Figure 11 (a), (b), and (c) depict PDE4 inhibitor compounds, (a) nitraquazone,
5 (b) denbufylline, and (c) rolipram useful in the present invention.

Figure 12 (a), (b), and (c) depict PDE4 inhibitor compounds, (a) GI198425X, (b) CP-77059, and (c) GI193600X, useful in the present invention.

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DETAILED DESCRIPTION OF THE INVENTION

As used herein, the term "alkyl" refers to a straight or branched chain hydrocarbon having from one to twelve carbon atoms, optionally substituted with substituents selected from the group which includes, but is not limited to, C₁-C₆ alkyl,
15 C₁-C₆ alkoxy, C₁-C₆ alkylsulfanyl, C₁-C₆ alkylsulfenyl, C₁-C₆ alkylsulfonyl, oxo, hydroxy, mercapto, amino optionally substituted by alkyl, carboxy, carbamoyl optionally substituted by alkyl, aminosulfonyl optionally substituted by a substituent selected from the group including alkyl, nitro, cyano, halogen and lower perfluoroalkyl, multiple degrees of substitution being allowed. Examples of "alkyl" as used herein
20 include, but are not limited to, methyl, ethyl, n-butyl, n-pentyl, isobutyl, isopropyl and the like.

As used herein, the term "alkoxy" refers to the group R_aO-, where R_a is alkyl.

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As used herein, the term "aryl" refers to a 6 to 10 carbon aromatic moiety. Examples of "aryl" as used herein include, but are not limited to, phenyl, pyridyl, pyrimidyl, pyridazyl, pyrazyl, and triazyl.

As used herein, the term "aryloxy" refers to the group R_aO-, where R_a is aryl.

30

As used herein the term "halogen" refers to the group fluorine, chlorine, bromine, and iodine.

As used herein the term "neoplasm" refers to an abnormal growth of cells or
5 tissue and is understood to include benign, i.e., non-cancerous growths, and
malignant, i.e., cancerous growths. The term "neoplastic" means of or related to a
neoplasm.

As used herein the term "agent" is understood to mean a substance that
10 produces a desired effect in a tissue, system, animal, mammal, human, or other
subject. Accordingly, the term "anti-neoplastic agent" is understood to mean a
substance producing an anti-neoplastic effect in a tissue, system, animal, mammal,
human, or other subject. It is also to be understood that an "agent" may be a single
compound or a combination or composition of two or more compounds.

15

As used herein, the term "solvate" is a complex of variable stoichiometry
formed by a solute and a solvent. Solvents, by way of example, include water,
methanol, ethanol, or acetic acid.

20 As used herein the term "progenitor cells" refers to parent or precursor cells
that are typically proliferating rapidly and give rise to cells that function in a tissue,
system, animal, mammal, human, or other subject. Such progenitor cells include, but
are not limited to, bone marrow stem cells that give rise to erythrocytes, leukocytes
and platelets; epithelial progenitors including epithelial stem cells of the intestinal
25 tract, and hair follicle progenitors which give rise to hair.

As used herein the term "specific stage progenitor" refers to
progenitors/precursors that can be assessed by a colony forming assay, e.g., CFU-GM
an early (immature) granulocyte macrophage progenitor. As used herein the term
30 "multiple stage progenitors" refers to a mixture of early and late (mature)

progenitors/ precursors that can still divide (not terminally differentiated) and produce daughter cells.

As used herein the term "phase specific anti-neoplastic compound" refers to anti-neoplastic compounds that are cell cycle specific. That is, the compounds act at or effect cells at specific phases of the cell cycle. For instance, there are about six dozen approved products for the treatment of malignancy. Many of the existing anti-neoplastic agents share a common mechanism of action and can be grouped together. These groups include alkylating agents such as melphalan, chlorambucil, cyclophosphamide, mechlorethamine, hexamethylmelamine, busulfan, carmustine, lomustine, and dacarbazine; antimetabolites such as 5-fluorouracil, methotrexate, cytarabine, mecaptopurine and thioguanine; antimitotic agents such as paclitaxel, vinblastine, vincristine; topoisomerase I inhibitors such as camptothecin and camptothecin derivatives; topoisomerase II inhibitors such as doxorubicin; and platinum coordination complexes such as cisplatin and carboplatin. These antineoplastic agents may either induce cytotoxic effects in a cell-cycle specific manner, i.e., are phase specific, or bind DNA and act in a non cell-cycle specific manner, i.e., are non-cell cycle specific (See Table 1).

As used herein, the term "effective amount" means that amount of a drug or pharmaceutical agent that will elicit the biological or medical response of a tissue, system, animal, mammal, human, or other subject that is being sought by a researcher or clinician. Furthermore the term "therapeutically effective amount" means any amount which, as compared to a corresponding subject who has not received such amount, results in improved treatment, healing, prevention, or amelioration of a disease, disorder, or side effect, or a decrease in the rate of advancement of a disease or disorder, and also includes amounts effective to enhance normal physiological function.

As recited above, in one embodiment a method of treating cancer is provided which includes administering a therapeutically effective amount of a PDE4 inhibitor to

a mammal. In another embodiment, a method of treating cancer is provided which includes administering therapeutically effective amounts of an anti-neoplastic agent that adversely affects normal progenitor cells upon use and a PDE4 inhibitor. Typically, any anti-neoplastic agent that adversely affects normal progenitor cells may
5 be utilized in the present invention. Such adverse affects include, but are not limited to myelosuppression, mucositis, and alopecia.

Of special interest, are phase specific anti-neoplastic agents. Typical phase specific anti-neoplastic agents useful in the present invention include, but are not
10 limited to, diterpenoids, vinca alkaloids, epipodophyllotoxins, antimetabolites, and camptothecins.

Diterpenoids, which are derived from natural sources, are phase specific anti - cancer agents that operate at the G₂/M phases of the cell cycle. It is believed that the
15 diterpenoids stabilize the β -tubulin subunit of the microtubules, by binding with this protein. Disassembly of the protein appears then to be inhibited with mitosis being arrested and cell death following. Examples of diterpenoids include, but are not limited to, paclitaxel and its analog docetaxel.

20 Paclitaxel, 5 β ,20-epoxy-1,2 α ,4,7 β ,10 β ,13 α -hexa-hydroxytax-11-en-9-one 4,10-diacetate 2-benzoate 13-ester with (2R,3S)-N-benzoyl-3-phenylisoserine; is a natural diterpene product isolated from the Pacific yew tree *Taxus brevifolia* and is commercially available as an injectable solution TAXOL®. It is a member of the taxane family of terpenes. It was first isolated in 1971 by Wani et al. J. Am. Chem. Soc.,
25 93:2325. 1971), who characterized its structure by chemical and X-ray crystallographic methods. One mechanism for its activity relates to paclitaxel's capacity to bind tubulin, thereby inhibiting cancer cell growth. Schiff et al., Proc. Natl Acad. Sci. USA, 77:1561-1565 (1980); Schiff et al., Nature, 277:665-667 (1979); Kumar, J. Biol. Chem, 256: 10435-10441 (1981). For a review of synthesis and anticancer activity of some
30 paclitaxel derivatives see: D. G. I. Kingston *et al.*, Studies in Organic Chemistry vol. 26,

entitled "New trends in Natural Products Chemistry 1986", Attaur-Rahman, P.W. Le Quesne, Eds. (Elsevier, Amsterdam, 1986) pp 219-235.

Paclitaxel has been approved for clinical use in the treatment of refractory
5 ovarian cancer in the United States (Markman et al., Yale Journal of Biology and
Medicine, 64:583, 1991; McGuire et al., Ann. Intern. Med., 111:273, 1989) and for the
treatment of breast cancer (Holmes et al., J. Nat. Cancer Inst., 83:1797, 1991.) It is a
potential candidate for treatment of neoplasms in the skin (Einzig et. al., Proc. Am.
Soc. Clin. Oncol., 20:46) and head and neck carcinomas (Forastire et. al., Sem. Oncol.,
10 20:56, 1990). The compound also shows potential for the treatment of polycystic
kidney disease (Woo et al., Nature, 368:750. 1994), lung cancer and malaria.
Treatment of patients with paclitaxel results in bone marrow suppression (multiple cell
lineages, Ignoff, R.J. et. al, Cancer Chemotherapy Pocket Guide, 1998) related to the
duration of dosing above a threshold concentration (50nM) (Kearns, C.M. et. al.,
15 Seminars in Oncology, 3(6) p.16-23, 1995).

Docetaxel, (2R,3S)- N-carboxy-3-phenylisoserine, N-*tert*-butyl ester, 13-ester
with 5 β -20-epoxy-1,2 α ,4,7 β ,10 β ,13 α -hexahydroxytax-11-en-9-one 4-acetate 2-
benzoate, trihydrate; is commercially available as an injectable solution as
20 TAXOTERE®. Docetaxel is indicated for the treatment of breast cancer. Docetaxel is a
semisynthetic derivative of paclitaxel *q.v.*, prepared using a natural precursor, 10-
deacetyl-baccatin III, extracted from the needle of the European Yew tree. The dose
limiting toxicity of docetaxel is neutropenia.

25 Vinca alkaloids are phase specific anti-neoplastic agents derived from the
periwinkle plant. Vinca alkaloids act at the M phase (mitosis) of the cell cycle by
binding specifically to tubulin. Consequently, the bound tubulin molecule is unable to
polymerize into microtubules. Mitosis is believed to be arrested in metaphase with cell
death following. Examples of vinca alkaloids include, but are not limited to,
30 vinblastine, vincristine, and vinorelbine.

Vinblastine, vincaleukoblastine sulfate, is commercially available as VELBAN® as an injectable solution. Although, it has possible indication as a second line therapy of various solid tumors, it is primarily indicated in the treatment of testicular cancer and various lymphomas including Hodgkin's Disease; lymphocytic and histiocytic lymphomas. Myelosuppression is the dose limiting side effect of vinblastine.

Vincristine, vincaleukoblastine, 22-oxo-, sulfate, is commercially available as ONCOVIN® as an injectable solution. Vincristine is indicated for the treatment of acute leukemias and has also found use in treatment regimens for Hodgkin's and non-Hodgkin's malignant lymphomas. Alopecia and neurologic effects are the most common side effect of vincristine and to a lesser extent myelosuppression and gastrointestinal mucositis effects occur.

Vinorelbine, 3',4'-didehydro -4'-deoxy-C'-norvincaleukoblastine [R-(R*,R*)-2,3-dihydroxybutanedioate (1:2)(salt)], commercially available as an injectable solution of vinorelbine tartrate (NAVELBINE®), is a semisynthetic vinca alkaloid. Vinorelbine is indicated as a single agent or in combination with other chemotherapeutic agents, such as cisplatin, in the treatment of various solid tumours, particularly non-small cell lung, advanced breast, and hormone refractory prostate cancers. Myelosuppression is the most common dose limiting side effect of vinorelbine.

Epipodophyllotoxins are phase specific anti-neoplastic agents derived from the mandrake plant. Epipodophyllotoxins typically affect cells in the S and G₂ phases of the cell cycle by forming a ternary complex with topoisomerase II and DNA causing DNA strand breaks. The strand breaks accumulate and cell death follows. Examples of epipodophyllotoxins include, but are not limited to, etoposide and teniposide.

Etoposide, 4'-demethyl-epipodophyllotoxin 9[4,6-O-(R)-ethylidene-β-D-glucopyranoside], is commercially available as an injectable solution or capsules as VePESID® and is commonly known as VP-16. Etoposide is indicated as a single agent or in combination with other chemotherapy agents in the treatment of testicular and

non-small cell lung cancers. Myelosuppression is the most common side effect of etoposide. The incidence of leucopenia tends to be more severe than thrombocytopenia.

5 Teniposide, 4'-demethyl-epipodophyllotoxin 9[4,6-O-(R)-thenylidene- β -D-glucopyranoside], is commercially available as an injectable solution as VUMON® and is commonly known as VM-26. Teniposide is indicated as a single agent or in combination with other chemotherapy agents in the treatment of acute leukemia in children. Myelosuppression is the most common dose limiting side effect of
10 teniposide. Teniposide can induce both leucopenia and thrombocytopenia.

Antimetabolite neoplastic agents are phase specific anti-neoplastic agents that act at S phase (DNA synthesis) of the cell cycle by inhibiting DNA synthesis or by inhibiting purine or pyrimidine base synthesis and thereby limiting DNA synthesis.
15 Consequently, S phase does not proceed and cell death follows. Examples of antimetabolite anti-neoplastic agents include, but are not limited to, fluorouracil, methotrexate, cytarabine, mecaptopurine and thioguanine.

5-fluorouracil, 5-fluoro-2,4- (1H,3H) pyrimidinedione, is commercially
20 available as fluorouracil. Adminsitration of 5-fluorouracil leads to inhibition of thymidylate synthesis and is also incorporated into both RNA and DNA. The result typically is cell death. 5-fluorouracil is indicated as a single agent or in combination with other chemotherapy agents in the treatment of carcinomas of the breast, colon, rectum, stomach and pancreas. Myelosuppression and mucositis are dose limiting side
25 effects of 5-fluorouracil. Other fluoropyrimidine analogs include 5-fluoro deoxyuridine (floxuridine) and 5-fluorodeoxyuridine monophosphate.

Cytarabine, 4-amino-1- β -D-arabinofuranosyl-2 (1H)-pyrimidinone, is commercially available as CYTOSAR-U® and is commonly known as Ara-C. It is
30 believed that cytarabine exhibits cell phase specificity at S-phase by inhibiting DNA chain elongation by terminal incorporation of cytarabine into the growing DNA chain.

Cytarabine is indicated as a single agent or in combination with other chemotherapy agents in the treatment of acute leukemia. Other cytidine analogs include 5-azacytidine and 2',2'-difluorodeoxycytidine (gemcitabine). Cytarabine induces leucopenia, thrombocytopenia, and mucositis.

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Mercaptopurine, 1,7-dihydro-6H-purine-6-thione monohydrate, is commercially available as PURINETHOL®. Mercaptopurine exhibits cell phase specificity at S-phase by inhibiting DNA synthesis by an as of yet unspecified mechanism. Mercaptopurine is indicated as a single agent or in combination with
10 other chemotherapy agents in the treatment of acute leukemia. Myelosuppression and gastrointestinal mucositis are expected side effects of mercaptopurine at high doses. A useful mercaptopurine analog is azathioprine.

Thioguanine, 2-amino-1,7-dihydro-6H-purine-6-thione, is commercially
15 available as TABLOID®. Thioguanine exhibits cell phase specificity at S-phase by inhibiting DNA synthesis by an as of yet unspecified mechanism. Thioguanine is indicated as a single agent or in combination with other chemotherapy agents in the treatment of acute leukemia. Myelosuppression, including leucopenia, thrombocytopenia, and anemia, is the most common dose limiting side effect of
20 thioguanine administration. However, gastrointestinal side effects occur and can be dose limiting. Other purine analogs include pentostatin, erythrohydroxynonyladenine, fludarabine phosphate, and cladribine.

Methotrexate, N-[4[[[2,4-diamino-6-pteridiny] methyl]methylamino] benzoyl]-L-glutamic acid, is commercially available as methotrexate sodium.
25 Methotrexate exhibits cell phase effects specifically at S-phase by inhibiting DNA synthesis, repair and/or replication through the inhibition of dihydrofolic acid reductase which is required for synthesis of purine nucleotides and thymidylate. Methotrexate is indicated as a single agent or in combination with other
30 chemotherapy agents in the treatment of choriocarcinoma, meningeal leukemia, non-Hodgkin's lymphoma, and carcinomas of the breast, head, neck, ovary and bladder.

Myelosuppression (leucopenia, thrombocytopenia, and anemia) and mucositis are expected side effect of methotrexate administration.

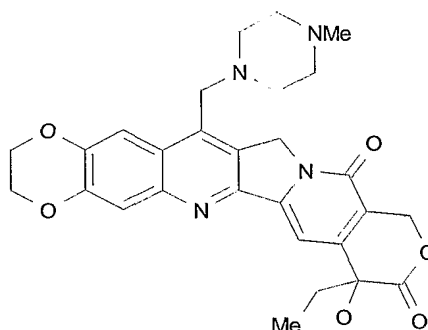
Camptothecins, including, camptothecin and camptothecin derivatives are
5 Topoisomerase I inhibitors. Camptothecins cytotoxic activity is believed to be related to its Topoisomerase I inhibitory activity. Examples of camptothecins include, but are not limited to topotecan, irinotecan and the various optical forms of 7-(4-methylpiperazino-methylene)-10,11-ethylenedioxy-20-camptothecin.

10 Topotecan HCl, (S)-10-[(dimethylamino)methyl]-4-ethyl-4,9-dihydroxy-1H-pyrano[3',4',6,7]indolizino[1,2-b]quinoline-3,14(4H,12H)-dione monohydrochloride, is commercially available as the injectable solution HYCAMTIN®. Topotecan is a semi-synthetic derivative of camptothecin which binds to the topoisomerase I – DNA complex and prevents religation of single strand breaks in DNA caused by
15 topoisomerase I. It is believed that cytotoxicity occurs as a result of irreparable double strand breaks caused by interaction of the topoisomerase I : DNA : topotecan ternary complex with replication enzymes. Topotecan is indicated for the treatment of metastatic carcinoma of the ovary and small cell lung cancer. The dose limiting side effect of topotecan HCl is myelosuppression, primarily neutropenia.

20

Irinotecan HCl, (4S)-4,11-diethyl-4-hydroxy-9-[(4-piperidinopiperidino)carbonyloxy]-1H-pyrano[3',4',6,7]indolizino[1,2-b]quinoline-3,14(4H,12H)-dione hydrochloride, is commercially available as the injectable solution CAMPTOSAR®. Irinotecan is a derivative of camptothecin which binds, along with its active
25 metabolite SN-38, to the topoisomerase I – DNA complex. It is believed that cytotoxicity occurs as a result of irreparable double strand breaks caused by interaction of the topoisomerase I : DNA : irinotecan or SN-38 ternary complex with replication enzymes. Irinotecan is indicated for treatment of metastatic cancer of the colon or rectum. The dose limiting side effects of irinotecan HCl are
30 myelosuppression, including neutropenia, and GI effects, including diarrhea.

Also of interest, is the camptothecin derivative of formula II following, currently under development, including the racemic mixture (R,S) form as well as the R and S enantiomers:



II

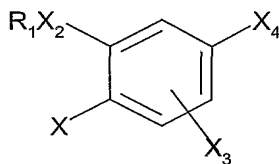
known by the chemical name "7-(4-methylpiperazino-methylene)-10,11-ethylenedioxy-20(R,S)-camptothecin (racemic mixture) or "7-(4-methylpiperazino-methylene)-10,11-ethylenedioxy-20(R)-camptothecin (R enantiomer) or "7-(4-methylpiperazino-methylene)-10,11-ethylenedioxy-20(S)-camptothecin (S enantiomer). Such compound as well as related compounds are described, including methods of making, in U.S. Patent Nos. 6,063,923; 5,342,947; 5,559,235; 5,491,237 and pending U.S. patent Application No. 08/977,217 filed November 24, 1997.

In one embodiment, the anti-neoplastic agent that adversely affects normal progenitor cells is a phase specific anti-neoplastic agent. In one embodiment, the phase specific anti-neoplastic agent includes a compound selected from diterpenoids, vinca alkaloids, epipodophyllotoxins, antimetabolites, camptothecins and pharmaceutically acceptable salts thereof. In another embodiment, the phase specific anti-neoplastic agent is a diterpenoid or pharmaceutically acceptable salt thereof. In a preferred embodiment, the phase specific anti-neoplastic agent is paclitaxel. In an alternative preferred embodiment, the phase specific anti-neoplastic agent is topotecan.

As recited above, the method of the present invention also includes administering a specific PDE4 inhibitor agent. Generally, any pharmaceutical agent having specific PDE4 inhibitor activity may be utilized in the present invention.

5 Typical PDE4 inhibitor agents include, but are not limited to, rolipram and rolipram derivatives; and fused ring compounds such as benzopyrazoles, benzimidazoles, benzofurans, diazepino-indoles, quinolines, quinolones, nitraquazone derivatives, purines and xanthines, and losartan analogues. Such PDE4 inhibitors are described, for instance, in "PDE4 inhibitors 1998", Norman, Peter; Exp. Opin. Ther. Patents (1998)
 10 8(7); and "Chronic Pulmonary Inflammation and Other Therapeutic Applications of PDEIV Inhibitors", Stafford, Jeffrey A. and Feldman, Paul L., Ch. 8 Annual Reports In Medicinal Chemistry Ch. 8, 71-80 (1996) which references are herein incorporated by reference to the extent of their disclosure of PDE4 inhibitor compounds.

15 One class of PDE4 inhibitor compounds that may be usefully employed in the present invention includes compounds of the Formula I :



20

wherein: R_1 is $-(CR_4 R_5)_n C(O)O(CR_4 R_5)_m R_6$, $-(CR_4 R_5)_n C(O)NR_4 (CR_4 R_5)_m R_6$, $-(CR_4 R_5)_n O(CR_4 R_5)_m R_6$, or $--(CR_4 R_5)_r R_6$ wherein the alkyl moieties may be optionally substituted with one or more halogens;

25 m is 0 to 2;

n is 0 to 4;

r is 0 to 6;

R_4 and R_5 are independently selected from hydrogen or a C_{1-2} alkyl;

R_6 is hydrogen, methyl, hydroxyl, aryl, halo substituted aryl, aryloxy C_{1-3} alkyl, halo substituted aryloxy C_{1-3} alkyl, indanyl, indenyl, C_{7-11} polycycloalkyl, tetrahydrofuranyl, furanyl, tetrahydropyranyl, pyranal, tetrahydrothienyl, thienyl, tetrahydrothiopyranyl, thiopyranyl, C_{3-6} cycloalkyl, or a C_{4-6} cycloalkyl containing one or two unsaturated bonds, wherein the cycloalkyl and heterocyclic moieties may be optionally substituted by 1 to 3 methyl groups or one ethyl group;

provided that:

when R_6 is hydroxyl, then m is 2; or

10 when R_6 is hydroxyl, then r is 2 to 6; or

when R_6 is 2-tetrahydropyranyl, 2-tetrahydrothiopyranyl, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then m is 1 or 2; or

when R_6 is 2-tetrahydropyranyl, 2-tetrahydrothiopyranyl, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then r is 1 to 6;

15 when n is 1 and m is 0, then R_6 is other than H in $--(CR_4 R_5)_n O (CR_4 R_5)_m$

R_6 ;

X is YR_2 , halogen, nitro, $NR_4 R_5$, or formyl amine;

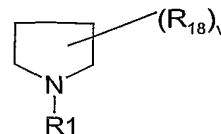
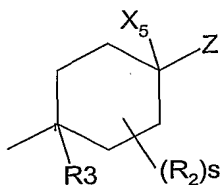
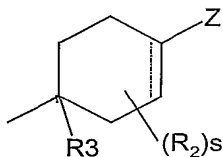
Y is O or $S(O)_{m'}$;

m' is 0, 1, or 2;

20 X_2 is O or NR_8 ;

X_3 is hydrogen or X;

X_4 is



25

$C(Z')NHR_{15}$, $CR_4(NOC(O)R_{16})$, or $CR_2'R_3'CR_4'R_5'R_6'$;

- X_5 is H, R_9 , OR_8 , CN, $C(O)R_8$, $C(O)OR_8$, $C(O)NR_8R_8$, or NR_8R_8 ;
 R_2 is independently selected from the group consisting of $--CH_3$ and $--CH_2CH_3$ optionally substituted by 1 or more halogens;
- 5 R_2' is a hydrogen, halogen, or OR_{17} ;
 s is 0 to 4;
 R_3' and R_4' are each independently $-(CH_2)_tX_6$;
 t is 0, 1, 2, or 3;
 v is 0, 1, 2, or 3;
- 10 X_6 is a mono- or bicyclic aryl group optionally containing one or more heteroatoms selected from sulfur, oxygen, or nitrogen;
 R_5' and R_6' are each independently hydrogen or an optionally substituted alkyl;
 R_3 is hydrogen, halogen, C_{1-4} alkyl, $CH_2NHC(O)C(O)NH_2$, halo-substituted C_{1-4} alkyl, $CH=CR_8R_8'$, cyclopropyl optionally substituted by R_8' , CN, OR_8 , CH_2OR_8 , NR_8R_{10} ,
- 15 $CH_2NR_8R_{10}$, $C(Z')H$, $C(O)OR_8$, $C(O)OR_8$, $C(O)NR_8R_{10}$, or $C\equiv CR_8$;
 Z' is O, S, NR_8 , NOR_8 , NCN , $C(-CN)_2$, CR_8CN , CR_8NO_2 , $CR_8C(O)OR_8$, $CR_8C(O)ONR_8R_8$, $C(-CN)NO_2$, $C(-CN)C(O)OR_9$, $C(-CN)C(O)NR_8R_8$;
 Z is $C(Y')R_{14}$, $C(O)OR_{14}$, $C(Y')NR_{10}R_{14}$, $C(NR_{10})NR_{10}R_{14}$, CN, $C(NOR_8)R_{14}$, $C(O)NR_8NR_8$, $C(O)R_8$, $C(O)NR_8NR_{10}R_{14}$, $C(NOR_{14})R_8$, $C(NR_8)NR_{10}R_{14}$, $C(NR_{14})NR_8R_8$, $C(NCN)NR_{10}R_{14}$,
- 20 $C(NCN)SR_9$, (2-, 4- or 5-imidazolyl), (3-, 4- or 5-pyrazolyl), (4- or 5-triazolyl[1,2,3]), (3- or 5-triazolyl[1,2,4]), (5-tetrazolyl), (2-, 4- or 5-oxazolyl), (3-, 4- or 5-isoxazolyl), (3- or 5-oxadiazolyl[1,2,4]), (2-oxadiazolyl[1,3,4]), (2-thiadiazolyl[1,3,4]), (2-, 4-, or 5-thiazolyl), (2-, 4-, or 5-oxazolidinyl), (2-, 4-, or 5-thiazolidinyl), or (2-, 4-, or 5-imidazolidinyl); wherein all of the heterocyclic ring systems may be optionally
- 25 substituted one or more times by R_{14} ;
 the dotted line in the first formula of X_4 represents a single or double bond;
 Y' is O or S;
- R_7 is $-(CR_4R_5)_qR_{12}$ or C_{1-6} alkyl wherein the R_{12} or C_{1-6} alkyl group is optionally substituted one or more times by C_{1-2} alkyl optionally substituted by one to three
- 30 fluorines, $--F$, $--Br$, $--Cl$, $--NO_2$, $--NR_{10}R_{11}$, $--C(O)R_8$, $--C(O)OR_8$, $--OR_8$, $--CN$, $--C(O)NR_{10}R_{11}$, $--OC(O)NR_{10}R_{11}$, $--OC(O)R_8$, $--NR_{10}C(O)NR_{10}R_{11}$, $--NR_{10}C(O)R_{11}$, $--NR_{10}C$

(O)OR₉, --NR₁₀ C(O)R₁₃, --C(NR₁₀)NR₁₀ R₁₁, --C(NCN)NR₁₀ R₁₁, --C(NCN)SR₉, --NR₁₀ C(NCN)SR₉, --NR₁₀ C(NCN)NR₁₀ R₁₁, --NR₁₀ S(O)₂ R₉, --S(O)_m R₉, --NR₁₀ C(O)C(O)NR₁₀ R₁₁, --NR₁₀ C(O)C(O)R₁₀, thiazolyl, imidazolyl, oxazolyl, pyrazolyl, triazolyl, or tetrazolyl;

q is 0, 1, or 2;

- 5 R₁₂ is C₃-C₇-cycloalkyl, (2-, 3- or 4-pyridyl), pyrimidyl, pyrazolyl, (1- or 2-imidazolyl), thiazolyl, triazolyl, pyrrolyl, piperazinyl, piperidinyl, morpholinyl, furanyl, (2- or 3-thienyl), (4- or 5-thiazolyl), quinolinyl, naphthyl, or phenyl;

R₈ is independently selected from hydrogen or R₉ ;

R₈ ' is R₈ or fluorine;

- 10 R₉ is C₁₋₄ alkyl optionally substituted by one to three fluorines;

R₁₀ is OR₈ or R₁₁ ;

R₁₁ is hydrogen, or C₁₋₄ alkyl optionally substituted by one to three fluorines; or when R₁₀ and R₁₁ are as NR₁₀ R₁₁ they may together with the nitrogen form a 5 to 7 membered ring optionally containing at least one additional heteroatom selected from

- 15 O/N/or S;

R₁₃ is oxazolidinyl, oxazolyl, thiazolyl, pyrazolyl, triazolyl, tetrazolyl, imidazolyl, imidazolidinyl, thiazolidinyl, isoxazolyl, oxadiazolyl, or thiadiazolyl, and each of these heterocyclic rings is connected through a carbon atom and each may be unsubstituted or substituted by one or two C₁₋₂ alkyl groups;

- 20 R₁₄ is hydrogen or R₇ ; or when R₁₀ and R₁₄ are as NR₁₀ R₁₄ they may together with the nitrogen form a 5 to 7 membered ring optionally containing one or more additional heteroatoms selected from O, N, or S;

R₁₅ is optionally substituted pyridyl, pyrazinyl, pyrimidinyl, isoxazolyl; or an N-oxide thereof; or a optionally substituted phenyl, wherein s optional substituents are one or
25 more selected from the group consisting of halogen, haloalkyl, aryl, arylalkyl, alkoxy, aryloxy, alkylthio, arylthio, alkoxycarbonyl, alkanoyl, aroyl, alkylsulphonyl, arylsulphonyl, alkylsulphinyl, arylsulphinyl, hydroxy, hydroxyalkyl, formyl, alkanoylamino, aroylamino, cyano, and nitro;

R₁₆ is amino, C₁-C₆ alkylamino, arylamino, C₁-C₆ alkoxy, or aryloxy;

- 30 R₁₇ is hydrogen, or optionally substituted alkyl, alkenyl, alkoxyalkyl, or alkanoyl, or a formyl, carboxamido, or thiocarboxamido;

R_{18} is (=O), hydrogen, $-C(O)R_2$, $-CH_3$, $-CH_2CH_3$, $-C(O)OH$;

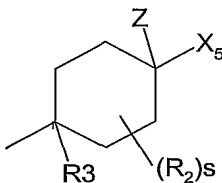
provided that:

when R_{12} is N-pyrazolyl, N-imidazolyl, N-triazolyl, N-pyrrolyl, N-piperazinyl, N-piperidinyl, or N-morpholinyl, then q is not 1; or

5 the pharmaceutically acceptable salts thereof.

In one embodiment, X_2 is O; R_1 is $(CR_4R_5)_rR_6$, where $r=0$ and R_6 is C_3 - C_6 cycloalkyl, preferably cyclopentyl; $X=YR_2$; R_2 is $-CH_3$; $Y=O$;

X_4 is



10

R_3 is $-CN$; X_5 is hydrogen; $s=0$; Z is $C(O)OR_{14}$; and R_{14} is hydrogen.

15

Of particular interest of the compounds of Formula I is the PDE4 inhibitor *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof. This compound has also been referred to as ariflo and is depicted in Figure 9(a). The compound and related compounds as well as the synthesis of such compounds is described in U.S. Patent No. 5,643,946, said patent being incorporated herein by reference to extent that it teaches PDE4 inhibitor compounds and synthesis of the same.

20

Another group of PDE4 inhibitor compounds of Formula I that may be usefully employed in the present invention includes piclamilast, i.e., N-(3,5-dichloropyrid-4-yl)-3-cyclopentyloxy-4-methoxybenzamide, depicted in Figure 9(b), and related compounds. Piclamilast and related compounds are described, as well as the synthesis thereof, in International Patent Application WO92/12961 and WO 97/48697.

25

A further group of PDE4 inhibitor compounds from Formula I that can be usefully employed in the present invention includes filaminast, 1-[3-cyclopentyloxy]-4-methoxyphenyl] ethanone (E)-O-{aminocarbonyl}oxime, which is depicted in Figure 10(a), and related compounds. Filaminast and related compounds are described, as well as the synthesis thereof, in European Patent Application EP 470805A1.

Still another group of Formula I PDE4 inhibitor compounds that can be usefully employed in the process of the present invention includes (+/-)-4-[2-(3-cyclopentyloxy-4-methoxyphenyl)-2-phenylethyl]pyridine (CDP-840), which is depicted in Figure 10(b), and related compounds. These compounds are described, as well as the synthesis thereof, in International Patent Applications WO 94/14742; 97/23460; 97/23461; 97/38976; 97/42172; and 98/12178.

Additional PDE4 inhibitors which can be utilized in the present invention include, but are not limited to:

- (i) nitroquazone, depicted in Figure 11(a), and nitraquazone derivatives, such compounds being described in WO 93/07146 including the synthesis thereof;
- (ii) denbufylline, i.e., 7-acetonyl,1,3dibutylxanthine, made by SmithKline Beecham and depicted in Figure 11(b);
- (iii) rolipram, depicted in Figure 11(c);
- (iv) RS-25344, depicted in Figure 12(a) and related compounds, such compounds being described in WO 93/07146 and 93/19068 including the synthesis thereof;
- (v) CP-77059, depicted in Figure 12(b) and related compounds, such compounds being described in WO 96/40636 and 97/05105 including the synthesis thereof; and
- (vi) GI 193600X, available from Glaxo Wellcome, Inc., depicted in Figure 12(c).

In one embodiment, the PDE4 inhibitor agent is a compound of Formula I, In another embodiment, the PDE4 inhibitor is *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-

methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof; N-(3,5-dichloropyrid-4-yl)-3-cyclopentyloxy-4-methoxybenzamide; 1-[3-cyclopentyloxy)-4-methoxyphenyl] ethanone (E)-O-{aminocarbonyl}oxime; or (+/-)-4-[2-(3-cyclopentyloxy-4-methoxyphenyl)-2-phenylethyl]pyridine.

5

Rolipram, a PDE4 inhibitor agent, when used alone had little protective effect in the cell based assays utilized herein. This is in agreement with the lack of effect of rolipram when used alone on various neutrophil functions (Au B-T et al. British J Pharmacol, 123: 1260-1266, (1998); Nielsen et al. J Allergy Clin Immunol, 86: 801-808, (1990)). However, a significant synergism between PDE4 inhibitors and agents that activate adenylate cyclase and raise intracellular levels of cAMP, e.g. forskolin was observed (Nicholson, CD; Shahid, M. Pulm Pharmacol, 7: 1-17 (1994); Au B-T et al. British J Pharmacol, 123: 1260-1266, (1998); Nielsen et al. J Allergy Clin Immunol, 86: 801-808, (1990)). While not wishing to be bound by theory, the present inventors believe that in human cancer patients an activator of adenylate cyclase may not be needed because serum levels of the adenylate cyclase activator prostaglandin E2 (PGE₂), in these patients are elevated (Young MR, Cancer Met. Rev., 13: 337-348 (1994); Uotila P., Cancer Immunol Immunother, 43: 1-9, (1996); Baxevanis CN et al. Eur J Cancer, 33 (8): 1202-1208, (1997)), thus contributing to elevated cAMP levels.

20

However, in an alternative embodiment, the method of treating cancer of the present invention further includes administering a therapeutically effective amount of an agent effective in raising intracellular concentrations of cAMP or analogs thereof along with the phase specific cancer agent and the PDE4 inhibitor agent. Suitable compounds include agents which (1) increase cAMP levels by activating surface receptors which are G_s protein coupled to the cAMP generating enzyme adenylate cyclase¹ including, but not limited to, prostaglandin E2, β_2 -adrenoreceptor agonists, adenosine A1 receptor agonists, and adenosine A2 receptor agonists; and (2) increase cAMP levels by directly stimulating adenylate cyclase, including, but not limited to forskolin, and stable analogs of cAMP including, but not limited to, dibutyl cAMP; 8-bromo c-AMP. Furthermore, it is also to be understood that the agent, which is

30

effective to raise intracellular cAMP concentrations, may be at least one additional PDE4 inhibitor.

In one embodiment, the agent, which is effective to raise intracellular levels of cAMP, is a β -2 adrenoreceptor agonist. Typical examples include, but are not limited to, terbutaline, albuterol, pirbuterol, bitolterol, metaproterenol, formoterol, and salmeterol or salt thereof.

Terbutaline, (+/-)- α [(tert-butylamino)methyl]-3,5-dihydroxybenzyl alcohol sulfate (2:1 salt), is commercially available as BRETHINE®. Terbutaline is described in U.S. Patent 3,938,838. Albuterol, racemic α^1 [(tert-butylamino) methyl]-4-hydroxy-m-xylene- α , α' -diol sulfate (2:1 salt), is commercially available as PROVENTIL®. Pirbuterol, (R,S) α^6 -[[[(1,1-dimethylethyl)amino]methyl]-3-hydroxy-2,6-pyridinedimethanolmonoacetate salt, is commercially available as MAXAIR™. Bitolterol, 4-[2-[(1,1-dimethylethyl)amino]-1-hydroxyethyl]-1,2-phenylene-4-methylbenzoate methanesulfonate salt, is commercially available as TORNALATE®. Metaproterenol, 1-(3,5-dihydroxyphenyl)-2-isopropylaminoethanol sulfate, is commercially available as ALUPENT®. Metaproterenol is described in U.S. patent 3,341,594. Formoterol, [(R*,R*)-(+/-)-N-[2-hydroxy-5-[1-hydroxy-2[[2-(4-methoxyphenyl)-1-methylethyl]ethyl]phenyl]formamide], is described in U.S. patent 3,994,974.

In a preferred embodiment, the β -2 adrenoreceptor agonist is salmeterol or salt thereof. Salmeterol, 4-hydroxy- α^1 [[[(6-(4-phenylbutoxy)hexyl)amino) methyl]-1,3-benzenedimethanol, preferably the 1-hydroxy-2-naphthalenecarboxylate salt (xinasoate), is commercially available as SEREVENT® and is described in GB 2140800.

In another embodiment, the phase specific anti-neoplastic agent is paclitaxel, the PDE4 inhibitor is *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof and the agent effective to raise intracellular levels of cAMP is salmeterol or salt thereof.

In an alternative embodiment, the phase specific anti-neoplastic agent is topotecan, the PDE4 inhibitor is *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl] cyclohexane-1-carboxylic acid or salt thereof and the agent effective to raise
5 intracellular levels of cAMP is salmeterol or salt thereof.

The animal requiring treatment using the method of the present invention is a mammal, such as a human being.

10 The anti-neoplastic agent and the PDE4 inhibitor may be employed in combination in accordance with the invention by administration concomitantly in (1) a unitary pharmaceutical composition including both compounds or (2) separate pharmaceutical compositions each including one of the compounds. Alternatively, the combination may be administered separately in a sequential manner wherein the PDE4
15 inhibitor or anti-neo-plastic agent is administered first and the other second. Such sequential administration may be close in time or remote in time. In a presently preferred embodiment, the PDE4 inhibitor is administered first and the anti-neoplastic agent is subsequently administered within a time period suitable to achieve the desired therapeutic effect. When utilized, in one embodiment, the agent, which is
20 effective to raise intracellular cAMP concentrations is administered concomitantly with the PDE4 inhibitor compound and before or after administration of the anti-neoplastic agent. In another embodiment, the agent, which is effective to raise intracellular cAMP concentrations is administered concomitantly with the PDE4 inhibitor compound and the anti-neoplastic agent. In a further embodiment, the
25 agent which is effective to raise intracellular cAMP concentrations is administration sequentially with the phase specific anti-neoplastic agent and the PDE4 inhibitor agent in any sequence and within any time period that is suitable to achieve the desired therapeutic effect. In all applications of the method of the present invention the PDE4 inhibitor agent and if utilized the agent which is effective to raise
30 intracellular cAMP concentrations are administered in a manner with the phase specific anti-neoplastic agent, so as to achieve a preventative or ameliorative effect

on side effects that would otherwise be presented in the absence of the PDE4 inhibitor agent and if utilized the agent which is effective to raise intracellular cAMP concentrations.

5 The anti-neoplastic agents, PDE4 inhibitor agents and agents which are effective to raise intracellular cAMP concentrations of the present invention may be administered by any appropriate route. Suitable routes include oral, rectal, nasal, topical (including buccal and sublingual), vaginal, and parenteral (including subcutaneous, intramuscular, intravenous, intradermal, intrathecal, and epidural). It
10 will be appreciated that the preferred route may vary with, for example, the condition of the recipient of the combination. It will also be appreciated that each of the agents/activators administered may be administered by the same or different routes and that one or more of the phase specific anti-neoplastic agent, PDE4 inhibition agent and adenylate cyclase inhibitor may be compounded together in a
15 pharmaceutical composition/formulation.

 The method of the present invention may also be employed with other therapeutic methods of cancer treatment. In particular, in anti-neoplastic therapy, combination therapy with other chemotherapeutic, hormonal, antibody agents as well
20 as surgical and/or radiation treatments other than those mentioned above are envisaged. Combination therapies according to the present invention thus include the administration of at least one anti-neoplastic agent and at least one PDE4 inhibitor agent and optionally at least one agent which is effective to raise intracellular cAMP concentrations as well optional use of other therapeutic agents including other anti-
25 neoplastic agents. Such combination of agents may be administered together or separately and, when administered separately this may occur simultaneously or sequentially in any order, both close and remote in time. The amounts of the compound(s) of formula (I) and the other pharmaceutically active agent(s) and the relative timings of administration will be selected in order to achieve the desired
30 combined therapeutic effect.

 Pharmaceutical formulations adapted for oral administration may be presented as discrete units such as capsules or tablets; powders or granules; solutions or

suspensions in aqueous or non-aqueous liquids; edible foams or whips; or oil-in-water liquid emulsions or water-in-oil liquid emulsions.

For instance, for oral administration in the form of a tablet or capsule, the
5 active drug component can be combined with an oral, non-toxic pharmaceutically acceptable inert carrier such as ethanol, glycerol, water and the like. Powders are prepared by comminuting the compound to a suitable fine size and mixing with a similarly comminuted pharmaceutical carrier such as an edible carbohydrate, as, for example, starch or mannitol. Flavoring, preservative, dispersing and coloring agent can
10 also be present.

Capsules are made by preparing a powder mixture as described above, and filling formed gelatin sheaths. Glidants and lubricants such as colloidal silica, talc, magnesium stearate, calcium stearate or solid polyethylene glycol can be added to the
15 powder mixture before the filling operation. A disintegrating or solubilizing agent such as agar-agar, calcium carbonate or sodium carbonate can also be added to improve the availability of the medicament when the capsule is ingested.

Moreover, when desired or necessary, suitable binders, lubricants,
20 disintegrating agents and coloring agents can also be incorporated into the mixture. Suitable binders include starch, gelatin, natural sugars such as glucose or beta-lactose, corn sweeteners, natural and synthetic gums such as acacia, tragacanth or sodium alginate, carboxymethylcellulose, polyethylene glycol, waxes and the like. Lubricants used in these dosage forms include sodium oleate, sodium stearate, magnesium
25 stearate, sodium benzoate, sodium acetate, sodium chloride and the like. Disintegrators include, without limitation, starch, methyl cellulose, agar, bentonite, xanthan gum and the like. Tablets are formulated, for example, by preparing a powder mixture, granulating or slugging, adding a lubricant and disintegrant and pressing into tablets. A powder mixture is prepared by mixing the compound, suitably comminuted,
30 with a diluent or base as described above, and optionally, with a binder such as carboxymethylcellulose, an aliginate, gelatin, or polyvinyl pyrrolidone, a solution retardant such as paraffin, a resorption accelerator such as a quaternary salt and/or an absorption agent such as bentonite, kaolin or dicalcium phosphate. The powder mixture can be granulated by wetting with a binder such as syrup, starch paste, acadia

mucilage or solutions of cellulosic or polymeric materials and forcing through a screen. As an alternative to granulating, the powder mixture can be run through the tablet machine and the result is imperfectly formed slugs broken into granules. The granules can be lubricated to prevent sticking to the tablet forming dies by means of
5 the addition of stearic acid, a stearate salt, talc or mineral oil. The lubricated mixture is then compressed into tablets. The compounds of the present invention can also be combined with free flowing inert carrier and compressed into tablets directly without going through the granulating or slugging steps. A clear or opaque protective coating consisting of a sealing coat of shellac, a coating of sugar or polymeric material and a
10 polish coating of wax can be provided. Dyestuffs can be added to these coatings to distinguish different unit dosages.

Oral fluids such as solution, syrups and elixirs can be prepared in dosage unit form so that a given quantity contains a predetermined amount of the compound.
15 Syrups can be prepared by dissolving the compound in a suitably flavored aqueous solution, while elixirs are prepared through the use of a non-toxic alcoholic vehicle. Suspensions can be formulated by dispersing the compound in a non-toxic vehicle. Solubilizers and emulsifiers such as ethoxylated isostearyl alcohols and polyoxy ethylene sorbitol ethers, preservatives, flavor additive such as peppermint oil or
20 natural sweeteners or saccharin or other artificial sweeteners, and the like can also be added.

Where appropriate, dosage unit formulations for oral administration can be microencapsulated. The formulation can also be prepared to prolong or sustain the
25 release as for example by coating or embedding particulate material in polymers, wax or the like.

The agents for use according to the present invention can also be administered in the form of liposome delivery systems, such as small unilamellar vesicles, large
30 unilamellar vesicles and multilamellar vesicles. Liposomes can be formed from a variety of phospholipids, such as cholesterol, stearylamine or phosphatidylcholines.

Agents for use according to the present invention may also be delivered by the use of monoclonal antibodies as individual carriers to which the compound molecules
35 are coupled. The compounds may also be coupled with soluble polymers as targetable

drug carriers. Such polymers can include polyvinylpyrrolidone, pyran copolymer, polyhydroxypropylmethacrylamide-phenol, polyhydroxyethylaspartamidephenol, or polyethyleneoxidepolylysine substituted with palmitoyl residues. Furthermore, the compounds may be coupled to a class of biodegradable polymers useful in achieving controlled release of a drug, for example, polylactic acid, polepsilon caprolactone, polyhydroxy butyric acid, polyorthoesters, polyacetals, polydihdropyrans, polycyanoacrylates and cross-linked or amphipathic block copolymers of hydrogels.

Pharmaceutical formulations adapted for transdermal administration may be presented as discrete patches intended to remain in intimate contact with the epidermis of the recipient for a prolonged period of time. For example, the active ingredient may be delivered from the patch by iontophoresis as generally described in Pharmaceutical Research, 3(6), 318 (1986).

Pharmaceutical formulations adapted for topical administration may be formulated as ointments, creams, suspensions, lotions, powders, solutions, pastes, gels, sprays, aerosols or oils.

For treatments of the eye or other external tissues, for example mouth and skin, the formulations are preferably applied as a topical ointment or cream. When formulated in an ointment, the active ingredient may be employed with either a paraffinic or a water-miscible ointment base. Alternatively, the active ingredient may be formulated in a cream with an oil-in-water cream base or a water-in-oil base.

Pharmaceutical formulations adapted for topical administrations to the eye include eye drops wherein the active ingredient is dissolved or suspended in a suitable carrier, especially an aqueous solvent.

Pharmaceutical formulations adapted for topical administration in the mouth include lozenges, pastilles and mouth washes.

Pharmaceutical formulations adapted for rectal administration may be presented as suppositories or as enemas.

Pharmaceutical formulations adapted for nasal administration wherein the carrier is a solid include a coarse powder having a particle size for example in the range 20 to 500 microns which is administered in the manner in which snuff is taken, i.e. by rapid inhalation through the nasal passage from a container of the powder held close up to the nose. Suitable formulations wherein the carrier is a liquid, for administration as a nasal spray or as nasal drops, include aqueous or oil solutions of the active ingredient.

Pharmaceutical formulations adapted for administration by inhalation include fine particle dusts or mists that may be generated by means of various types of metered dose pressurised aerosols, nebulizers or insufflators.

Pharmaceutical formulations adapted for vaginal administration may be presented as pessaries, tampons, creams, gels, pastes, foams or spray formulations.

Pharmaceutical formulations adapted for parenteral administration include aqueous and non-aqueous sterile injection solutions which may contain anti-oxidants, buffers, bacteriostats and solutes which render the formulation isotonic with the blood of the intended recipient; and aqueous and non-aqueous sterile suspensions which may include suspending agents and thickening agents. The formulations may be presented in unit-dose or multi-dose containers, for example sealed ampoules and vials, and may be stored in a freeze-dried (lyophilized) condition requiring only the addition of the sterile liquid carrier, for example water for injections, immediately prior to use. Extemporaneous injection solutions and suspensions may be prepared from sterile powders, granules and tablets.

It should be understood that in addition to the ingredients particularly mentioned above, the formulations may include other agents conventional in the art having regard to the type of formulation in question, for example those suitable for oral administration may include flavoring agents.

Salts encompassed within the term "pharmaceutically acceptable salts" refer to non-toxic salts of the compounds of this invention which are generally prepared by reacting a free base with a suitable organic or inorganic acid or by reacting the acid with a suitable organic or inorganic base.

Also, contemplated in the present invention is a pharmaceutical combination including an anti-neoplastic agent and a PDE4 inhibitor. In another embodiment, the pharmaceutical combination includes an anti-neoplastic agent, a PDE4 inhibitor, and
5 an agent, which is effective to raise intracellular cAMP concentrations. The anti-neoplastic agent, PDE4 inhibitor, and agent that is effective to raise intracellular cAMP concentrations are as described above. In another embodiment, the phase specific anti-neoplastic agent is paclitaxel, the PDE4 inhibitor is *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof and
10 if included the agent effective to raise intracellular levels of cAMP is salmeterol or salt thereof. In still another embodiment, the phase specific anti-neoplastic agent is topotecan, the PDE4 inhibitor is *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof and if included the agent effective to raise intracellular levels of cAMP is salmeterol or salt thereof.

15

The present invention provides a method for treating cancer whereby detrimental side effects associated with anti-neoplastic agents that adversely affect normal progenitor cells are attenuated. It will be understood by those skilled in the art that the specific cancers treated will depend on the specific anti-neoplastic agents
20 used. As such the expected side effects as well may vary depending on the dose, routes, and cycles of anti-neoplastic agents utilized to treat a specific tumor. Typical, side effects which may be attenuated include but are not limited to myelosuppression including neutropenia, thrombocytopenia, or anemia; mucositis; and alopecia. In a preferred embodiment, the side effect attenuated is myelosuppression or mucositis. In
25 a more preferred embodiment, the side effect attenuated is neutropenia.

As indicated, therapeutically effective amounts of the specific anti-neoplastic agent, the PDE4 inhibitor, and if utilized the agent, that is effective to raise intracellular cAMP concentrations, are administered to a mammal. Typically, the
30 therapeutically effective amount of one of the administered agents of the present invention will depend upon a number of factors including, for example, the age and

weight of the mammal, the precise condition requiring treatment, the severity of the condition, the nature of the formulation, and the route of administration. Ultimately, the therapeutically effective amount will be at the discretion of the attendant physician or veterinarian.

5

Typically, the anti-neoplastic agent will be given in the range of 0.1 to 100 mg/kg body weight of recipient (mammal) per day and more usually in the range of 1 to 10 mg/kg body weight per day. Acceptable daily dosages of the PDE4 inhibitor for preventing/reducing the severity of side effects induced by administration of a phase
10 specific anti-neoplastic agent, may be from about 0.1 to about 1000 mg/day, and preferably from about 0.2 to about 100 mg/day. Acceptable daily dosages of the agent which is effective to raise intracellular cAMP concentrations for stimulating cAMP production to assist in reducing the severity of side effects induced by administration of a anti-neoplastic agent, may also be from about 0.1 to about 1000 mg/day, and
15 preferably from about 0.2 to about 100 mg/day.

Also, provided in the present invention is use of (i) an anti-neoplastic agent which has an adverse effect on normal progenitor cells upon use; and (ii) a PDE4 inhibitor in the treatment of cancer. The anti-neoplastic agent and PDE4 agent are as
20 described above. This aspect may further include use of an agent, which is effective to raise intracellular levels of cAMP, in conjunction with the anti-neoplastic agent and PDE4 inhibitor. Such agent is as described above.

In addition, there is provided herein use of (i) an anti-neoplastic agent which
25 has an adverse effect on normal progenitor cells upon use; and (ii) a PDE4 inhibitor in the preparation of a medicament for the treatment of cancer. This aspect may further use an agent that is effective to raise intracellular levels of cAMP in conjunction with the anti-neoplastic agent and the PDE4 inhibitor. The anti-neoplastic agent, PDE4 inhibitor, and the agent that is effective to raise intracellular levels of cAMP are as
30 described above.

Also provided herein, is use of a PDE4 inhibitor in the preparation of a medicament for the attenuation of cancer treatment side effects on normal progenitor cells. The PDE4 inhibitor is as described above.

5

The following examples are intended for illustration only and are not intended to limit the scope of the invention in any way. The following abbreviations are used in the examples following as well as in other portions of the specification.

10	cAMP	cyclic adenosine monophosphate
	CELISA	cell-based enzyme linked immunoabsorbent assay
	CFU-GM	colony forming unit-granulocyte macrophages
	DMEM	Dulbecco's Modified Eagle's Media
	DMSO	dimethyl sulfoxide
15	ELISA	enzyme linked immunoabsorbent assay
	EDTA	ethylenediaminetetraacetic acid
	FACS	fluorescent-activated cell analysis
	FBS	fetal bovine serum
	G-CSF	granulocyte-colony stimulating factor
20	GM-CSF	granulocyte-macrophage-colony stimulating factor
	HBSS	Hank's Balanced Salt Solution
	IMDM	Iscoe's Modified Dulbecco's medium
	PBS	phosphate buffered saline
	PDE4	phosphodiesterase 4
25	RT	room temperature
	SRB	sulforhodamine B

Example 1

Preparation of PDE4 Stock Solutions.

30

PDE4 inhibitors were synthesized utilizing published procedures. The PDE4 inhibitors were prepared as 1.0 mM stock solutions in DMSO. The PDE4 inhibitors prepared were Rolipram (Baures, PW, 1993 J.Med. Chem. 36(3274-3277.); Ariflo (SB-207499, Christensen S.B. 1998 *J Med Chem* 41, 821-835); GI193600X (Stafford JA et al., Journal of Medicinal Chemistry, 38(26): 4972-4975, (1995)); RP-73401 (RPR Patent, WO9748697); CDP-840 (Merck, Celltech patents WO9723460, WO9723461, WO9738976, WO9742172, WO9812178); nitraquazone and denbufylline (Norman P., 1998, Exp. Opin. Ther. Patents, 8 (7)), 771-784); GI198425X (Syntex patents, WO-9307146, WO-9319068); and GI147404X (Pfizer Inc., WO-9640636, WO-9705105). The PDE4 stock solutions were utilized in the following examples.

Prevention of myelosuppression was shown by protection of neutrophil progenitors by PDE4 inhibitors from the cytotoxic effects of cancer chemotherapeutic agents. In Examples 2-7 a variety of chemically distinct PDE4 inhibitors were tested for their ability to protect neutrophil progenitors. Three methods, more completely described in the following examples, were utilized for a determination of cell survival and ability to produce daughter cells of primary human neutrophil progenitors: (1) a FACS assay which determined the percent of live cells by size and granularity; (2) a colony forming assay which measured the number of CFU-GM progenitors (a specific early stage neutrophil progenitor); and (3) a cell-based ELISA which measured CD15 antigen the value of which was proportional to the number of neutrophil lineage cells. This assay allows evaluation of neutrophil progenitors (cells that can divide and produce daughter neutrophil-lineage cells) at multiple stages of differentiation.

Example 2

Flow cytometric evaluation of expanded CD34⁺ cells treated with rolipram plus forskolin followed by treatment with paclitaxel.

Human bone marrow samples were collected from presumed healthy human donors following informed consent through an agreement with Duke University Medical Center, Durham, North Carolina. Mononuclear cells were isolated using a

1.077 Ficoll-Paque gradient according to the published procedure (Dornsife, RE et al., Antimicrob Agents Chemother, 35: 322-328, (1991)) and were suspended in HBSS with 2% FBS at a concentration of 5×10^7 cells/ml. CD34⁺ stem cells were isolated by "StemSep™ CD34⁺ stem cell selection kit" by following the instructions from the manufacturer (Stem Cell Technologies). Cells were adjusted to 1×10^6 cells/ml in complete growth media and maintained in a humidified, 5% CO₂ incubator. The complete growth media was IMDM supplemented with 20% FBS, 1% penicillin-streptomycin antibiotics, the recombinant human cytokines (Biosource) Flt-3/Flk-2 ligand (50 ng/ml), stem cell factor (50 ng/ml), interleukin-3 (40ng/ml), and G-CSF (10 ng/ml). The cells were maintained at a density of between 0.5 and 3×10^5 cells/ml. The CD34⁺ stem cells were expanded in culture over an average range of 11 to 23 days. Five groups of expanded CD34⁺ cells were treated with rolipram (1μM), plus Forskolin (100 μM), an adenylate cyclase activator, for 24 hours and five groups of expanded CD34⁺ cells were treated with DMSO for 24 hours. After 24 hours, to the 5 sets of cells that were treated with DMSO and 5 groups of cells treated with rolipram plus forskolin, paclitaxel was added to give final concentrations of 0, 30, 100, 300, and 600 nM of paclitaxel respectively and incubated for an additional 24 hours. After the 48 hours of treatment, the cells were counted, washed and re-suspended in PBS with 2% FBS. A total of 1×10^6 cells were pelleted and then fixed with 0.5% paraformaldehyde in PBS + 2% FBS solution. The prepared cells were analyzed on a BD FACSsort flow cytometer. The results are depicted in Figure 2.

The expanded CD34⁺ stem cells treated with rolipram plus forskolin prior to paclitaxel treatment, in Example 2, were analyzed on a flow cytometer (FACS) for surviving and apoptotic cell fractions (Fig. 2). The pretreatment with rolipram plus forskolin significantly reduced the apoptotic fraction in paclitaxel treated cells when compared with the cells pretreated with only DMSO. Conversely, the surviving cell fraction was much higher in cells pretreated with the chemoprotective regimen (Fig. 2). The surviving cell fractions after the pretreatment increased from 45% to 86% 27% to 81%, 23% to 63%, and 18% to 59% at 30nM, 100nM, 300nM, and 600nM of paclitaxel, respectively.

A T-test analysis (One-way ANOVA, with application of either equal or unequal variance [difference in standard error of the mean (sem) < or > 3-fold, respectively) was performed to compare between treatment groups to determine if statistically significant differences existed (MS Excel, SAS JMP). The pretreatment data (rolipram plus forskolin) were determined to be statistically significantly different from DMSO controls ($p = 0.0062$), indicating that more cells survived.

It is important to note that the expanded CD34⁺ stem cells could contain, in addition to the progenitors of neutrophils, the progenitors of platelets, erythrocytes, and/or lymphocytes (B, T, and NK). The protection of the expanded CD34⁺ cells from the cytotoxic effects of paclitaxel by the rolipram plus forskolin pretreatment suggests that the PDE4 inhibitors can potentially protect cancer patients from multiple blood lineage hematopoietic toxicities (neutropenia, thrombocytopenia, and anemia).

Example 3

Protection of specific early neutrophil progenitors (CFU-GMs) as measured by colony forming assay – rolipram plus forskolin followed by treatment with paclitaxel, 5-fluorouracil, or etoposide.

Expanded CD34⁺ cells, prepared according to the procedure of Example 2, were isolated and pretreated with DMSO or rolipram plus forskolin as described in Example 2. After 24 hours, varying amounts of paclitaxel, 5-fluorouracil, and etoposide were added and incubated for an additional 24 hours. After the 48 hours of treatment, the cells were counted, washed twice with 4 ml of HBSS supplemented with 2% FBS, and resuspended in complete growth media. The cells were then plated in a semi solid growth medium for the growth of CFU-GMs following a published method with the exception that the G-CSF (10ng/ml) was included in place of GM-CSF and erythropoietin (Dornsife, RE et al., Antimicrob Agents Chemother, 35: 322-328, (1991); Dornsife RE; Averett DR., Antimicrob Agents Chemother, 40(2): 514-519,

(1996)). The CFU-GMs were counted after 12 to 14 days. The results are depicted in Figure 3.

5 Cancer patients are particularly predisposed to bacterial infection due to the fact that they are usually immunosuppressed by both the cancer and anti-neoplastic treatments. Patients with neutrophil counts of 500/ μ l have a substantially increased susceptibility for infection. Neutrophil counts of even less confers an escalating risk as the neutrophil count drops.

10 To examine the potential to protect specific early neutrophil progenitors (CFU-GM) from the cytotoxic effects of anti-neoplastic drugs a colony forming assay (Fig. 3) was used after the cells were exposed for 48 hours of treatments in the previously specified liquid culture in Example 3. After the expanded CD34⁺ cells were treated with varying concentrations of 5-fluorouracil or etoposide the number of CFU-GM
15 colonies decreased in a dose dependent manner with increasing amounts of the drug (Fig. 3(a) & 3(b)). In contrast, expanded CD34⁺ cells treated with paclitaxel did not exhibit a good dose response (Fig. 3(c)). However, a pretreatment of the expanded CD34⁺ cells with rolipram, and forskolin, increased the survival of CFU-GM progenitors, as the number of resultant colonies were increased to a statistically significant level
20 (95% confidence) in cells pretreated with the chemoprotective regimen prior to a treatment with 5-fluorouracil ($p = 0.014$), etoposide ($p = 0.046$), or paclitaxel ($p = 0.012$) when compared with cells pretreated with only DMSO (Fig. 3).

Example 4

25 *Protection of multiple stage neutrophil progenitors as measured by CD15 neutrophil antigen cell-based ELISA (CELISA) method – rolipram plus forskolin followed by treatment with paclitaxel, 5-fluorouracil, etoposide, vinblastine, vinorelbine, methotrexate, or topotecan.*

30 Expanded CD34⁺ stem cells were isolated as described in Example 2 and grown in complete growth media. The expanded CD34⁺ cells were treated with DMSO or

rolipram plus forskolin as previously described. After 24h, varying amounts of paclitaxel, 5-fluorouracil, etoposide, vinblastine, vinorelbine, methotrexate, or topotecan were added and incubated for an additional 24 hours. After the treatment, the cells were counted, washed twice with 4 ml of HBSS supplemented with 2% FBS, and resuspended in complete growth media. The expression of CD15 antigen was used as a measure of the number of neutrophil lineage cells expanded from surviving progenitors of multiple stages of differentiation. (modified version of assay described in Warren et al., 1995 Stem Cells 13:167-174).

10 The cells were plated at a set number into 96-well plates in 100 μ l volume per well of complete media. In some experiments, an additional 100 μ l of complete media was added to plates maintained for later time points before harvest. The plates were harvested at different time points by centrifuging at 450 G for 10 minutes at room temperature (RT) and the cells were washed several times in HBSS by centrifugation of the plates. The cells were removed by suspension twice in HBSS with 5 mM EDTA with 100 μ l per well and transfer of the total 200 μ l volume to capture plates (Costar DNA-bind, #2498) that had been pretreated with 1% glutaraldehyde and stored (Warren et al., 1995 Stem Cells 13:167-174).

20 For CELISA, the preservative solution (PBS, 0.1%BSA, 0.1% azide) was removed from the capture plate and the wells were washed with 100 μ l PBS. A mouse anti-human CD15 antibody (Biosource, Clone B-H8) 0.5 μ g/well, 100 μ l PBS with 5% goat sera) was added and incubated at RT for 60 min. The plates were washed twice with 300 μ l PBS/well and incubated with a secondary antibody (goat anti-mouse IgM-biotin, Sigma B-9265, 0.105 μ g/well, 100 μ l PBS with 5% goat sera) at RT for 60 min. After three to six 200 μ l/well washes, streptavidin-horseradish peroxidase (Pierce 21126, 0.1 μ g/well in a 1% solution of stable peroxide buffer, Pierce 34062) was added. The plates were incubated for ten minutes and washed three to six times with 200 μ l /well. For the final detection step, the chemiluminescent reagent Supersignal[®] Ultra (Pierce 34075) prepared just prior to use by mixing equal parts of the luminol enhancer and the stable peroxide was added to the plates (100 μ l per well) and gently

mixed for five minutes at RT. The plates were read on a Wallace Victor II machine for one second for chemiluminescence at 425 nm.

The data from the Wallac Victor were transferred to Excel for analysis. Averages of replicate wells from treated cells were calculated along with standard error of the mean (sem) of the counts minus the background signal from replicate wells with media only (no cells). The results are depicted in Figure 4.

The protection of neutrophil progenitors from the cytotoxic effects of anti-neoplastic agents by pretreatment with rolipram in combination with forskolin was further confirmed in Example 4 by a CELISA method (Fig. 4). The expression of CD15, an antigen expressed on stage specific CFU-GM progenitors as well as other granulocytic (neutrophil) lineage cells, by CELISA was used to confirm the statistically significant protection of multiple stages of neutrophil progenitors. This assay measures the recovery and expansion of CD15+ cells from neutrophil progenitors (those able to divide and produce CD15+ daughter cells). The CD15 signal from the cell cultures decreased with increasing concentrations of paclitaxel, 5-fluorouracil, etoposide, vinblastine, vinorelbine, methotrexate, or topotecan (Fig. 4A-4G). A pretreatment of the expanded CD34+ cells with rolipram plus forskolin increased the CD15 signal antigen to a statistically significant level (99% confidence) (paclitaxel $p = 1.3 \times 10^{-11}$, 5-FU $p = 4.8 \times 10^{-7}$, etoposide $p = 5.8 \times 10^{-6}$, vinblastine $p = 4.4 \times 10^{-3}$, vinorelbine $p = 1.5 \times 10^{-9}$, methotrexate $p = 1.2 \times 10^{-7}$, and topotecan $p = 9.8 \times 10^{-8}$), indicating that more progenitors survived the cytotoxic effects of the anti-neoplastic agents to divide and produce more CD15+ daughter cells versus progenitors pretreated with only DMSO (Fig. 4).

Example 5

Protection of multiple stage neutrophil progenitors as measured by CD15 neutrophil antigen-based CELISA method –PDE4 inhibitors (other than rolipram) plus forskolin followed by treatment with paclitaxel.

Neutrophil progenitors were protected by PDE4 inhibitors (a) Ariflo, or (b) Piclamilast, (c) CDP-840 [GR259653X], (d) CDP-840 [GR259654X], (e) Nitraquazone, (f) Denbufylline, (g) RS-25344, (h) CP-77059, and (i) GI193600X each in combination with forskolin from the cytotoxic effects of paclitaxel as measured by CELISA.

5 Experimental details of this experiment are the same as described under Example 4 except that another PDE4 inhibitor (1 μ M) was substituted for rolipram. The results are depicted in Figure 5.

Example 5 illustrates the protection of neutrophil progenitors from the cytotoxic effects of paclitaxel by a variety of PDE4 specific inhibitors each in combination with forskolin as measured by the CELISA method. Pretreatment of expanded CD34+ cells with the different PDE4 inhibitors each plus forskolin increased expression of CD15+ signal to a statistically significant level (Ariflo $p = 2.7 \times 10^{-9}$, Piclamilast $p = 8.5 \times 10^{-15}$, CDP-840 [GR259653X] $p = 1.9 \times 10^{-5}$, CDP-840 [GR259654X] $p = 6.5 \times 10^{-13}$, Nitraquazone $p = 1.0 \times 10^{-8}$, Denbufylline $p = 1.1 \times 10^{-9}$, RS-25344 $p = 2.2 \times 10^{-6}$, CP-77059 $p = 3.2 \times 10^{-7}$, and GI193600X $p = 3.4 \times 10^{-13}$), suggesting that more neutrophil progenitors survived the cytotoxic effects of the anti-neoplastic agent paclitaxel (Fig. 5(a)- Fig. 5(i)).

20 Example 6

Protection of neutrophil progenitors as measured by CD15 neutrophil antigen CELISA method –Ariflo plus forskolin followed by treatment with topotecan.

Neutrophil progenitors were protected by ariflo plus forskolin from the cytotoxic effects of topotecan as measured by CELISA. Experimental details of this experiment are the same as described under Example 4 except that ariflo (1 μ M) was substituted for rolipram and topotecan was used as the anti-neoplastic agent. The results are depicted in Figure 6.

Example 6 illustrates the protection of neutrophil progenitors from the cytotoxic effects of topotecan by ariflo plus forskolin as measured by the CELISA method. Pretreatment of expanded CD34+ cells with ariflo plus forskolin increased

expression of CD15+ signal to a statistically significant level ($p = 1.5 \times 10^{-5}$), suggesting that more neutrophil progenitors survived the cytotoxic effects of the anti-neoplastic agent topotecan (Fig. 6).

5

Example 7

Protection of neutrophil progenitors as measured by CD15 neutrophil antigen CELISA method –rolipram plus forskolin followed by treatment with anti-neoplastic agents listed in Table 1.

10

Protection of neutrophil progenitors from the cytotoxic effects of different anti-neoplastic agents, listed in table 1, by the PDE4 inhibitor, rolipram used in combination with forskolin was measured by CELISA method according to Example 4. The results are depicted in Table 1.

15

TABLE 1

	Drug	Protection?	Target/mechanism	Cell cycle
20	Paclitaxel	YES	microtubules, stabilizer	G2/M
	Vinblastine	YES	tubulin, destabilizer	G2/M
	Vinorelbine	YES	blocks polymerization of tubules	M
25	Etoposide	YES	stabilizes topoisomerase II enzyme	G2/M
	5-FU	YES	anti-metabolite, DNA syn inhibitor	S
30	methotrexate	YES	purine synthesis inhibitor	S, G2
	topotecan	YES	topoisomerase I inhibitor	G2/M, S
	Cisplatin	NO	binds to DNA, X-link	non-specific
35	Carboplatin	NO	binds to DNA, different lesions than cisplatin : non-specific	

Doxorubicin NO intercalates DNA, stabilize topo II enzyme: non-specific

5

Protection of neutrophil progenitors from the cytotoxic effects of different anti-neoplastic agents by the PDE4 inhibitor rolipram in combination with forskolin was measured by CELISA method. The results are summarized in Table 1. Rolipram and forskolin protected the neutrophil progenitors from the agents that act in a cell cycle specific manner. In contrast, no protection was observed from the agents that bind DNA and act in a non-cell cycle specific manner. For instance, Figure 7 depicts results for non-cell specific neoplastic agents cisplatin, doxorubicin, and carboplatin pretreated with rolipram plus forskolin. For the effects of carboplatin the pretreatment of rolipram plus forskolin of the cells compared to DMSO showed no significant difference ($p = 0.55$). For cisplatin ($p = 0.027$) and doxorubicin ($p = 0.0032$), however the pretreatment with rolipram plus forskolin not only did not protect but appeared to significantly increase the toxicity towards the neutrophil progenitors.

Potential protection from mucositis induced by anti-neoplastic agents by pretreatment with PDE4 inhibitors is illustrated in Example 8. Following toxic insult by an anti-neoplastic agent, initial tissue damage and inflammation results in the death of proliferating epithelial cells. Tissue atrophy and ulceration are involved in the late phases of mucositis. An in vitro model using mink lung epithelial cells was used to examine whether a chemoprotection regimen including a PDE4 inhibitor would be useful for preventing mucositis. It is believed that a PDE4 inhibitor could potentially prevent mucositis in two ways: 1) by protecting epithelial progenitor cells and 2) by producing an anti-tumor necrosis factor effect (anti-inflammatory).

30 Example 8

Protection of mink lung epithelial cells by different PDE4 inhibitors in combination with forskolin from the cytotoxic effects of paclitaxel (10 nM).

CCL64 mink lung epithelial cells (ATCC # CCL-64) were grown in DMEM supplemented with 10% FBS and 1% penicillin-streptomycin and seeded into 96-well plates. After overnight incubation at 37°C, cells were treated with 100 μ M forskolin in combination with each of the PDE4 inhibitors (1 μ M). After 24 hours of treatment, paclitaxel was added to wells at various concentrations up to 10 nM. Time zero control plates were harvested by staining with Sulforhodamine B (Skehan P. et. al., (1990), J. Nat'l Cancer Inst 82:1107-12.) After an additional 48 hours of incubation at 37°C, the plates were harvested and stained with sulforhodamine B. The absorbance of the samples was read at the optical density of 562 nm. The no-drug controls were set to 100% and values were normalized to time zero control. The results are depicted in Table II.

TABLE II

PDE4 Inhibitor	% Increase in Survival
Rolipram	110
Ariflo	110
GI193600X	120
Picamilast	130
CDP-840 ¹	125
Nitraquazone	115
CDP-840 ²	110
RS25344	105
Denbufylline	95
CP-77059	85

¹ GR259653X

² GR259654X

As indicated in Example 8, cells were pre-treated with forskolin plus various PDE4 inhibitors (1 μ M) and then challenged with paclitaxel at various concentrations.

The sulforhodamine B signal from the cells was expressed as percent of control (no-drug, vehicle only). An increase in signal, indicating protection, ranged from 85% to 130% with the cells pre-treated with forskolin plus a particular PDE4 inhibitor and followed by a paclitaxel challenge at 10nM.

5

It is extremely important that an agent that protects normal cells against toxicity induced by anti-neoplastic agents should not impact the anti-tumor effects of the agent. In example 9, human cell lines representing colon, lung, and breast tumor types were tested with the chemoprotection regimen of either rolipram plus forskolin or Ariflo plus forskolin to examine whether they would or would not be protected from the anti-neoplastic effects of the cancer drug paclitaxel.

10

Example 9

No effect on the anti-tumor activity with human tumor cell lines HT-29 (colon), RKO (colon), lines A549 (lung), and MDA468 (breast) treated with forskolin plus rolipram or forskolin plus Ariflo followed by treatment with paclitaxel.

15

The human tumor cell lines HT-29 (colon: ATCC# HTB 38), RKO (colon, ATCC# HTB 130), lines A549 (lung, ATCC# CCL 185), and MDA468 (breast, ATCC# HTB 132) were obtained from the American Type Culture Collection and maintained in DMEM containing 10% FBS. Tumor cells were grown at 37°C in a humidified atmosphere of 5% CO₂. Adherent tumor cells were seeded into 96 well plates and incubated for six hours to allow for attachment. After six hours the chemoprotectants forskolin (100 μM) plus rolipram (1 μM) or forskolin (100 μM) plus ariflo (1μM) were added. After overnight incubation, paclitaxel (100 nM) was added for 24 hours. Cells were then washed free of chemoprotectants and paclitaxel, placed in fresh media, and grown for 5 days. Growth was measured daily using sulforhodamine B (Skehan P. et. al., (1990), J. Nat'l Cancer Inst 82:1107-12). The results are depicted in Figure 8.

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In Figure 8 (a), (b), (c) and (d), two different colon cancer lines (RKO, HT-29) were tested with either rolipram or ariflo as the PDE4 inhibitor in the

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chemoprotection regimen (forskolin plus PDE4 inhibitor). The cells were then followed by a challenge by 100 nM paclitaxel. Cells receiving the chemoprotectant regimen versus those that were pre-treated with DMSO when challenged with paclitaxel did not effect the inhibition of tumor cell growth when challenged with paclitaxel. The
5 control tumor cells that were not exposed to paclitaxel quickly expanded over the 72 hours.

In Figure 8 (e), (f), (g), and (h) similar results were seen with a breast cancer cell line (MDA468, monitored for growth every other day for 10 days) and lung cancer
10 cell line (A549). While the lung cell line A549 was not completely inhibited by paclitaxel, this cell line also had no additional protection with the chemoprotectant regimen of forskolin plus rolipram or forskolin plus Ariflo. These data clearly show that the chemoprotectant regimen of PDE4 inhibitor and forskolin does not protect
tumor cells from the anti-neoplastic effects of paclitaxel.

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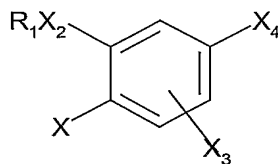
CLAIMS

We claim:

1. A method of treating cancer in a mammal, comprising: administering to said
5 mammal therapeutically effective amounts of (i) an anti-neoplastic agent which has an adverse effect on normal progenitor cells upon use; and (ii) a PDE4 inhibitor.
2. The method of claim 1, wherein the anti-neoplastic agent is a phase specific anti-neoplastic agent.
- 10 3. The method of claim 2, wherein the phase specific anti-neoplastic agent is selected from the group consisting of diterpenoids; vinca alkaloids; epipodo - phyllotoxins, antimetabolites, camptothecins, or pharmaceutically acceptable salts thereof.
- 15 4. The method of claim 1, wherein the anti-neoplastic agent is paclitaxel.
5. The method of claim 1, wherein the anti-neoplastic agent is topotecan.
6. The method of claim 1, wherein the PDE4 inhibitor is a compound of formula I

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I



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wherein: R_1 is $-(CR_4 R_5)_n C(O)O(CR_4 R_5)_m R_6$, $-(CR_4 R_5)_n C(O)NR_4 (CR_4 R_5)_m R_6$, $-(CR_4 R_5)_n O(CR_4 R_5)_m R_6$, or $--(CR_4 R_5)_r R_6$ wherein the alkyl moieties may be optionally substituted with one or more halogens;

m is 0 to 2;

5 n is 0 to 4;

r is 0 to 6;

R_4 and R_5 are independently selected from hydrogen or a C_{1-2} alkyl;

R_6 is hydrogen, methyl, hydroxyl, aryl, halo substituted aryl, aryloxy C_{1-3} alkyl, halo

substituted aryloxy C_{1-3} alkyl, indanyl, indenyl, C_{7-11} polycycloalkyl, tetrahydrofuranyl,

10 furanyl, tetrahydropyranyl, pyranal, tetrahydrothienyl, thienyl, tetrahydrothiopyranal,

thiopyranal, C_{3-6} cycloalkyl, or a C_{4-6} cycloalkyl containing one or two unsaturated

bonds, wherein the cycloalkyl and heterocyclic moieties may be optionally substituted

by 1 to 3 methyl groups or one ethyl group;

provided that:

15 when R_6 is hydroxyl, then m is 2; or

when R_6 is hydroxyl, then r is 2 to 6; or

when R_6 is 2-tetrahydropyranyl, 2-tetrahydrothiopyranal, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then m is 1 or 2; or

20 when R_6 is 2-tetrahydropyranyl, 2-tetrahydrothiopyranal, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then r is 1 to 6;

when n is 1 and m is 0, then R_6 is other than H in $--(CR_4 R_5)_n O(CR_4 R_5)_m R_6$

;

X is YR_2 , halogen, nitro, $NR_4 R_5$, or formyl amine;

Y is O or $S(O)_{m'}$;

25 m' is 0, 1, or 2;

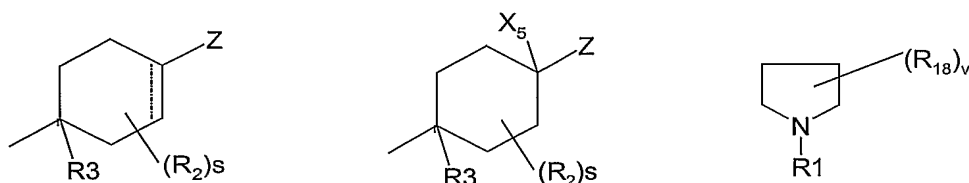
X_2 is O or NR_8 ;

X_3 is hydrogen or X;

X_4 is

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$C(Z')NHR_{15}$, $CR_4(NOC(O)R_{16})$, or $CR_2'R_3'CR_4'R_5'R_6'$;

- 10 X_5 is H, R_9 , OR_8 , CN, $C(O)R_8$, $C(O)OR_8$, $C(O)NR_8$, R_8 , or NR_8 R_8 ;
 R_2 is independently selected from the group consisting of $--CH_3$ and $--CH_2CH_3$
optionally substituted by 1 or more halogens;
 R_2' is a hydrogen, halogen, or OR_{17} ;
 s is 0 to 4;
- 15 R_3' and R_4' are each independently $-(CH_2)_tX_6$;
 t is 0, 1, 2, or 3;
 v is 0, 1, 2, or 3;
 X_6 is a mono- or bicyclic aryl group optionally containing one or more heteroatoms
selected from sulfur, oxygen, or nitrogen;
- 20 R_5' and R_6' are each independently hydrogen or an optionally substituted alkyl;
 R_3 is hydrogen, halogen, C_{1-4} alkyl, $-CH_2NHC(O)C(O)NH_2$, halo-substituted C_{1-4} alkyl, $-CH=CR_8'R_8'$, cyclopropyl optionally substituted by R_8' , CN, OR_8 , CH_2OR_8 , NR_8R_{10} ,
 $CH_2NR_8R_{10}$, $C(Z')H$, $C(O)OR_8$, $C(O)OR_8$, $C(O)NR_8R_{10}$, $-$, or $C\equiv CR_8$;
 Z' is O, S, NR_8 , NOR_8 , NCN , $C(-CN)_2$, CR_8CN , CR_8NO_2 , $CR_8C(O)OR_8$, $CR_8C(O)ONR_8R_8$, $C(-$
- 25 $CN)NO_2$, $C(-CN)C(O)OR_9$, $C(-CN)C(O)NR_8R_8$;
 Z is $C(Y')R_{14}$, $C(O)OR_{14}$, $C(Y')NR_{10}$ R_{14} , $C(NR_{10})NR_{10}$ R_{14} , CN, $C(NOR_8)R_{14}$, $C(O)NR_8$ NR_8 $C(O)R_8$,
 $C(O)NR_8$ NR_{10} R_{14} , $C(NOR_{14})R_8$, $C(NR_8)NR_{10}$ R_{14} , $C(NR_{14})NR_8$ R_8 $C(NCN)NR_{10}$ R_{14} , $C(NCN)SR_9$,

(2-, 4- or 5-imidazolyl), (3-, 4- or 5-pyrazolyl), (4- or 5-triazolyl[1,2,3]), (3- or 5-triazolyl[1,2,4]), (5-tetrazolyl), (2-, 4- or 5-oxazolyl), (3-, 4- or 5-isoxazolyl), (3- or 5-oxadiazolyl[1,2,4]), (2oxadiazolyl[1,3,4]), (2-thiadiazolyl[1,3,4]), (2-, 4-, or 5-thiazolyl), (2-, 4-, or 5-oxazolidinyl), (2-, 4-, or 5-thiazolidinyl), or (2-, 4-, or 5-imidazolidinyl);

5 wherein all of the heterocyclic ring systems may be optionally substituted one or more times by R_{14} ;

the dotted line in the first formula of X_4 represents a single or double bond;

Y' is O or S;

R_7 is $-(CR_4 R_5)_q R_{12}$ or C_{1-6} alkyl wherein the R_{12} or C_{1-6} alkyl group is optionally
 10 substituted one or more times by C_{1-2} alkyl optionally substituted by one to three fluorines, $--F$, $--Br$, $--Cl$, $--NO_2$, $--NR_{10} R_{11}$, $--C(O)R_8$, $--C(O)OR_8$, $--OR_8$, $--CN$, $--C(O)NR_{10} R_{11}$, $--OC(O)NR_{10} R_{11}$, $--OC(O)R_8$, $--NR_{10} C(O)NR_{10} R_{11}$, $--NR_{10} C(O)R_{11}$, $--NR_{10} C(O)OR_9$, $--NR_{10} C(O)R_{13}$, $--C(NR_{10})NR_{10} R_{11}$, $--C(NCN)NR_{10} R_{11}$, $--C(NCN)SR_9$, $--NR_{10} C(NCN)SR_9$, $--NR_{10} C(NCN)NR_{10} R_{11}$, $--NR_{10} S(O)_2 R_9$, $--S(O)_m R_9$, $--NR_{10} C(O)C(O)NR_{10} R_{11}$, $--NR_{10} C(O)C(O)R_{10}$, thiazolyl, imidazolyl, oxazolyl, pyrazolyl, triazolyl, or tetrazolyl;

q is 0, 1, or 2;

R_{12} is $C_3 -C_7$ -cycloalkyl, (2-, 3- or 4-pyridyl), pyrimidyl, pyrazolyl, (1- or 2-imidazolyl), thiazolyl, triazolyl, pyrrolyl, piperazinyl, piperidinyl, morpholinyl, furanyl, (2- or 3-thienyl), (4- or 5-thiazolyl), quinolinyl, naphthyl, or phenyl;

20 R_8 is independently selected from hydrogen or R_9 ;

R_8' is R_8 or fluorine;

R_9 is C_{1-4} alkyl optionally substituted by one to three fluorines;

R_{10} is OR_8 or R_{11} ;

R_{11} is hydrogen, or C_{1-4} alkyl optionally substituted by one to three fluorines; or when

25 R_{10} and R_{11} are as $NR_{10} R_{11}$ they may together with the nitrogen form a 5 to 7 membered ring optionally containing at least one additional heteroatom selected from O/N/or S;

R_{13} is oxazolidinyl, oxazolyl, thiazolyl, pyrazolyl, triazolyl, tetrazolyl, imidazolyl, imidazolidinyl, thiazolidinyl, isoxazolyl, oxadiazolyl, or thiadiazolyl, and each of these heterocyclic rings is connected through a carbon atom and each may be unsubstituted
 30 or substituted by one or two C_{1-2} alkyl groups;

R₁₄ is hydrogen or R₇ ; or when R₁₀ and R₁₄ are as NR₁₀ R₁₄ they may together with the nitrogen form a 5 to 7 membered ring optionally containing one or more additional heteroatoms selected from O, N, or S;

R₁₅ is optionally substituted pyridyl, pyrazinyl, pyrimidinyl, isoxazolyl; or an N-oxide thereof; or a optionally substituted phenyl, wherein s optional substituents are one or more selected from the group consisting of halogen, haloalkyl, aryl, arylalkyl, alkoxy, aryloxy, alkylthio, arylthio, alkoxy carbonyl, alkanoyl, aroyl, alkylsulphonyl, arylsulphonyl, alkylsulphinyl, arylsulphinyl, hydroxy, hydroxyalkyl, formyl, alkanoylamino, aroylamino, cyano, and nitro;

R₁₆ is amino, C₁-C₆ alkylamino, arylamino, C₁-C₆ alkoxy, or aryloxy;

R₁₇ is hydrogen, or optionally substituted alkyl, alkenyl, alkoxyalkyl, or alkanoyl, or a formyl, carboxamido, or thiocarboxamido;

R₁₈ is (=O), hydrogen, -C(O)R₂, -CH₃, -CH₂CH₃, -C(O)OH;

provided that:

when R₁₂ is N-pyrazolyl, N-imidazolyl, N-triazolyl, N-pyrrolyl, N-piperazinyl, N-piperidinyl, or N-morpholinyl, then q is not 1; or the pharmaceutically acceptable salts thereof.

7. The method of claim 1, wherein the PDE4 inhibitor is selected from the group consisting of

cis-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof;

N-(3,5-dichloropyrid-4-yl)-3-cyclopentyloxy-4-methoxybenzamide;

1-[3-cyclopentyloxy)-4-methoxyphenyl] ethanone (E)-O-

{aminocarbonyl} oxime; and

(+/-)-4-[2-(3-cyclopentyloxy-4-methoxyphenyl)-2-phenylethyl]pyridine.

8. The method of claim 1, wherein the PDE4 inhibitor is *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.

9. The method of claim 1, further comprising administering a therapeutically effective amount of an agent effective in raising intracellular concentrations of cAMP or analogs thereof.
- 5 10. The method of claim 9, wherein the agent is salmeterol or a salt thereof.
11. A method of treating cancer in a mammal, comprising: administering to said mammal therapeutically effective amounts of (i) paclitaxel; and (ii) *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.
- 10 12. The method of claim 11, further comprising administering a therapeutically effective amount of salmeterol or salt thereof.
13. A method of treating cancer in a mammal, comprising: administering to said
- 15 mammal therapeutically effective amounts of (i) topotecan; and (ii) *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.
14. The method of claim 13, further comprising administering a therapeutically effective amount of salmeterol or salt thereof.
- 20 15. A cancer treatment combination, comprising: therapeutically effective amounts of (i) an anti-neoplastic agent that adversely affects normal progenitor cells upon use; and (ii) a PDE4 inhibitor.
- 25 16. The combination of claim 15, wherein the anti-neoplastic agent is a phase specific anti-neoplastic agent.
17. The combination of claim 15, wherein the phase specific anti-neoplastic agent is selected from the group consisting of diterpenoids; vinca alkaloids; epipodo -
- 30 phyllotoxins, antimetabolites, camptothecins, or pharmaceutically acceptable salts thereof.

18. The combination of claim 15, wherein the anti-neoplastic agent is paclitaxel.

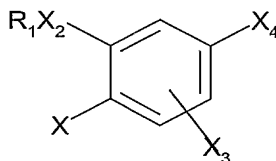
19. The combination of claim 15, wherein the anti-neoplastic agent is topotecan.

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20. The combination of claim 15, wherein the PDE4 inhibitor is a compound of formula I

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I



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wherein: R_1 is $-(CR_4 R_5)_n C(O)O(CR_4 R_5)_m R_6$, $-(CR_4 R_5)_n C(O)NR_4 (CR_4 R_5)_m R_6$, $-(CR_4 R_5)_n O(CR_4 R_5)_m R_6$, or $-(CR_4 R_5)_r R_6$ wherein the alkyl moieties may be optionally substituted with one or more halogens;

20 m is 0 to 2;

n is 0 to 4;

r is 0 to 6;

R_4 and R_5 are independently selected from hydrogen or a C_{1-2} alkyl;

R_6 is hydrogen, methyl, hydroxyl, aryl, halo substituted aryl, aryloxy C_{1-3} alkyl, halo

25 substituted aryloxy C_{1-3} alkyl, indanyl, indenyl, C_{7-11} polycycloalkyl, tetrahydrofuranyl, furanyl, tetrahydropyranyl, pyranal, tetrahydrothienyl, thienyl, tetrahydrothiopyranal, thiopyranal, C_{3-6} cycloalkyl, or a C_{4-6} cycloalkyl containing one or two unsaturated

bonds, wherein the cycloalkyl and heterocyclic moieties may be optionally substituted by 1 to 3 methyl groups or one ethyl group;

provided that:

- when R_6 is hydroxyl, then m is 2; or
 5 when R_6 is hydroxyl, then r is 2 to 6; or
 when R_6 is 2-tetrahydropyranyl, 2-tetrahydrothiopyranyl, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then m is 1 or 2; or
 when R_6 is 2-tetrahydropyranyl, 2-tetrahydrothiopyranyl, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then r is 1 to 6;
 10 when n is 1 and m is 0, then R_6 is other than H in $--(CR_4 R_5)_n O(CR_4 R_5)_m R_6$

;

X is YR_2 , halogen, nitro, $NR_4 R_5$, or formyl amine;

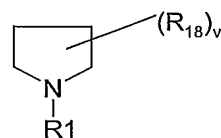
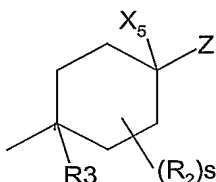
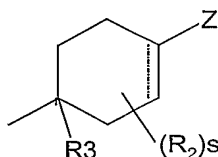
Y is O or $S(O)_{m'}$;

m' is 0, 1, or 2;

- 15 X_2 is O or NR_8 ;

X_3 is hydrogen or X;

X_4 is



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$C(Z')NHR_{15}$, $CR_4(NOC(O)R_{16})$, or $CR_2'R_3'CR_4'R_5'R_6'$;

- 25 X_5 is H, R_9 , OR_8 , CN, $C(O)R_8$, $C(O)OR_8$, $C(O)NR_8 R_8$, or $NR_8 R_8$;

R₂ is independently selected from the group consisting of --CH₃ and --CH₂ CH₃ optionally substituted by 1 or more halogens;

R₂' is a hydrogen, halogen, or OR₁₇;

s is 0 to 4;

5 R₃' and R₄' are each independently -(CH₂)_tX₆;

t is 0, 1, 2, or 3;

v is 0, 1, 2, or 3;

X₆ is a mono- or bicyclic aryl group optionally containing one or more heteroatoms selected from sulfur, oxygen, or nitrogen;

10 R₅' and R₆' are each independently hydrogen or an optionally substituted alkyl;

R₃ is hydrogen, halogen, C₁₋₄ alkyl, , CH₂NHC(O)C(O)NH₂, halo-substituted C₁₋₄ alkyl, -CH=CR₈'R₈', cyclopropyl optionally substituted by R₈ ', CN, OR₈, CH₂OR₈, NR₈R₁₀, CH₂NR₈R₁₀, C(Z')H, C(O)OR₈, C(O)OR₈, C(O)NR₈R₁₀, , or C≡CR₈;

Z' is O, S, NR₈, NOR₈, NCN, C(-CN)₂, CR₈CN, CR₈NO₂, CR₈C(O)OR₈, CR₈C(O)ONR₈R₈, C(-

15 CN)NO₂, C(-CN)C(O)OR₉, C(-CN)C(O)NR₈R₈;

Z is C(Y')R₁₄, C(O)OR₁₄, C(Y')NR₁₀ R₁₄, C(NR₁₀)NR₁₀ R₁₄, CN, C(NOR₈)R₁₄, C(O)NR₈ NR₈ C(O)R₈, C(O)NR₈ NR₁₀ R₁₄, C(NOR₁₄)R₈, C(NR₈)NR₁₀ R₁₄, C(NR₁₄)NR₈ R₈ C(NCN)NR₁₀ R₁₄, C(NCN)SR₉, (2-, 4- or 5-imidazolyl), (3-, 4- or 5-pyrazolyl), (4- or 5-triazolyl[1,2,3]), (3- or 5-triazolyl[1,2,4]), (5-tetrazolyl), (2-, 4- or 5-oxazolyl), (3-, 4- or 5-isoxazolyl), (3- or 5-oxadiazolyl[1,2,4]), (2oxadiazolyl[1,3,4]), (2-thiadiazolyl[1,3,4]), (2-, 4-, or 5-thiazolyl), (2-, 4-, or 5-oxazolidinyl), (2-, 4-, or 5-thiazolidinyl), or (2-, 4-, or 5-imidazolidinyl); wherein all of the heterocyclic ring systems may be optionally substituted one or more times by R₁₄ ;

the dotted line in the first formula of X₄ represents a single or double bond;

25 Y' is O or S;

R₇ is --(CR₄ R₅)_q R₁₂ or C₁₋₆ alkyl wherein the R₁₂ or C₁₋₆ alkyl group is optionally substituted one or more times by C₁₋₂ alkyl optionally substituted by one to three fluorines, --F, --Br, --Cl, --NO₂, --NR₁₀ R₁₁, --C(O)R₈, --C(O)OR₈, --OR₈, --CN, --C(O)NR₁₀ R₁₁, --OC(O)NR₁₀ R₁₁, --OC(O)R₈, --NR₁₀ C(O)NR₁₀ R₁₁, --NR₁₀ C(O)R₁₁, --NR₁₀ C(O)OR₉, --NR₁₀ C(O)R₁₃, --C(NR₁₀)NR₁₀ R₁₁, --C(NCN)NR₁₀ R₁₁, --C(NCN)SR₉, --NR₁₀ C(NCN)SR₉, --

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NR₁₀ C(NCN)NR₁₀ R₁₁, --NR₁₀ S(O)₂ R₉, --S(O)_m 'R₉, --NR₁₀ C(O)C(O)NR₁₀ R₁₁, --NR₁₀ C(O)C(O)R₁₀, thiazolyl, imidazolyl, oxazolyl, pyrazolyl, triazolyl, or tetrazolyl;

q is 0, 1, or 2;

- 5 R₁₂ is C₃ -C₇ -cycloalkyl, (2-, 3- or 4-pyridyl), pyrimidyl, pyrazolyl, (1- or 2-imidazolyl), thiazolyl, triazolyl, pyrrolyl, piperazinyl, piperidinyl, morpholinyl, furanyl, (2- or 3-thienyl), (4- or 5-thiazolyl), quinolinyl, naphthyl, or phenyl;

R₈ is independently selected from hydrogen or R₉ ;

R₈ ' is R₈ or fluorine;

R₉ is C₁₋₄ alkyl optionally substituted by one to three fluorines;

- 10 R₁₀ is OR₈ or R₁₁ ;

R₁₁ is hydrogen, or C₁₋₄ alkyl optionally substituted by one to three fluorines; or when R₁₀ and R₁₁ are as NR₁₀ R₁₁ they may together with the nitrogen form a 5 to 7 membered ring optionally containing at least one additional heteroatom selected from O/N/or S;

- 15 R₁₃ is oxazolidinyl, oxazolyl, thiazolyl, pyrazolyl, triazolyl, tetrazolyl, imidazolyl, imidazolidinyl, thiazolidinyl, isoxazolyl, oxadiazolyl, or thiadiazolyl, and each of these heterocyclic rings is connected through a carbon atom and each may be unsubstituted or substituted by one or two C₁₋₂ alkyl groups;

- 20 R₁₄ is hydrogen or R₇ ; or when R₁₀ and R₁₄ are as NR₁₀ R₁₄ they may together with the nitrogen form a 5 to 7 membered ring optionally containing one or more additional heteroatoms selected from O, N, or S;

- 25 R₁₅ is optionally substituted pyridyl, pyrazinyl, pyrimidinyl, isoxazolyl; or an N-oxide thereof; or a optionally substituted phenyl, wherein s optional substituents are one or more selected from the group consisting of halogen, haloalkyl, aryl, arylalkyl, alkoxy, aryloxy, alkylthio, arylthio, alkoxycarbonyl, alkanoyl, aroyl, alkylsulphonyl, arylsulphonyl, alkylsulphinyl, arylsulphinyl, hydroxy, hydroxyalkyl, formyl, alkanoylamino, aroylamino, cyano, and nitro;

R₁₆ is amino, C₁-C₆ alkylamino, arylamino, C₁-C₆ alkoxy, or aryloxy;

R₁₇ is hydrogen, or optionally substituted alkyl, alkenyl, alkoxyalkyl, or alkanoyl, or a formyl, carboxamido, or thiocarboxamido;

- 30 R₁₈ is (=O), hydrogen, -C(O)R₂, -CH₃, -CH₂CH₃, -C(O)OH; provided that:

when R₁₂ is N-pyrazolyl, N-imidazolyl, N-triazolyl, N-pyrrolyl, N-piperazinyl, N-piperidinyl, or N-morpholinyl, then q is not 1; or the pharmaceutically acceptable salts thereof.

- 5 21. The combination of claim 15, wherein the PDE4 inhibitor is selected from the group consisting of
- cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof;
- N-(3,5-dichloropyrid-4-yl)-3-cyclopentyloxy-4-methoxybenzamide;
- 10 1-[3-cyclopentyloxy)-4-methoxyphenyl] ethanone (E)-O-{aminocarbonyl} oxime; and
- (+/-)-4-[2-(3-cyclopentyloxy-4-methoxyphenyl)-2-phenylethyl]pyridine.
22. The combination of claim 15, wherein the PDE4 inhibitor is *cis*-4-cyano-4-[3-
- 15 (cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.
23. The combination of claim 15, further comprising (iii) a therapeutically effective amount of an agent effective in raising intracellular concentrations of cAMP or analogs thereof.
- 20 24. The combination of claim 23 wherein the agent is salmeterol or salt thereof.
25. A cancer treatment combination, comprising: therapeutically effective amounts of (i) paclitaxel; and (ii) *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-
- 25 methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.
26. The combination of claim 25, further comprising a therapeutically effective amount of salmeterol or salt thereof.

27. A cancer treatment combination, comprising: therapeutically effective amounts of (i) topotecan; and (ii) *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.

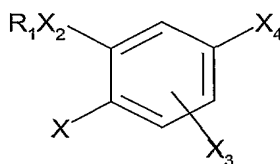
5 28. The combination of claim 27, further comprising a therapeutically effective amount of salmeterol or salt thereof.

29. A method of attenuating, in a mammal, cancer treatment side effects on normal
10 progenitor cells in the mammal, comprising: administering to the mammal a therapeutically effective amount of a PDE4 inhibitor.

30. The method of claim 29, wherein the PDE4 inhibitor is a compound of formula I

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I



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wherein: R_1 is $-(CR_4 R_5)_n C(O)O(CR_4 R_5)_m R_6$, $-(CR_4 R_5)_n C(O)NR_4 (CR_4 R_5)_m R_6$, $-(CR_4 R_5)_n O(CR_4 R_5)_m R_6$, or $--(CR_4 R_5)_r R_6$ wherein the alkyl moieties may be optionally substituted with one or more halogens;

m is 0 to 2;

25 n is 0 to 4;

r is 0 to 6;

R_4 and R_5 are independently selected from hydrogen or a C_{1-2} alkyl;

R₆ is hydrogen, methyl, hydroxyl, aryl, halo substituted aryl, aryloxy C₁₋₃ alkyl, halo substituted aryloxy C₁₋₃ alkyl, indanyl, indenyl, C₇₋₁₁ polycycloalkyl, tetrahydrofuranyl, furanyl, tetrahydropyranyl, pyranal, tetrahydrothienyl, thienyl, tetrahydrothiopyranyl, thiopyranyl, C₃₋₆ cycloalkyl, or a C₄₋₆ cycloalkyl containing one or two unsaturated bonds, wherein the cycloalkyl and heterocyclic moieties may be optionally substituted by 1 to 3 methyl groups or one ethyl group;

provided that:

when R₆ is hydroxyl, then m is 2; or

when R₆ is hydroxyl, then r is 2 to 6; or

when R₆ is 2-tetrahydropyranyl, 2-tetrahydrothiopyranyl, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then m is 1 or 2; or

when R₆ is 2-tetrahydropyranyl, 2-tetrahydrothiopyranyl, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then r is 1 to 6;

when n is 1 and m is 0, then R₆ is other than H in $-(\text{CR}_4 \text{ R}_5)_n \text{O}(\text{CR}_4 \text{ R}_5)_m \text{R}_6$

;

X is YR₂, halogen, nitro, NR₄ R₅, or formyl amine;

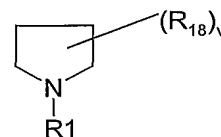
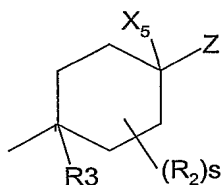
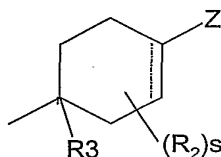
Y is O or S(O)_{m'};

m' is 0, 1, or 2;

X₂ is O or NR₈;

X₃ is hydrogen or X;

X₄ is



$C(Z')NHR_{15}$, $CR_4(NOC(O)R_{16})$, or $CR_2'R_3'CR_4'R_5'R_6'$;

X_5 is H, R_9 , OR_8 , CN, $C(O)R_8$, $C(O)OR_8$, $C(O)NR_8$, R_8 , or NR_8 , R_8 ;

R_2 is independently selected from the group consisting of $--CH_3$ and $--CH_2CH_3$

5 optionally substituted by 1 or more halogens;

R_2' is a hydrogen, halogen, or OR_{17} ;

s is 0 to 4;

R_3' and R_4' are each independently $-(CH_2)_tX_6$;

t is 0, 1, 2, or 3;

10 v is 0, 1, 2, or 3;

X_6 is a mono- or bicyclic aryl group optionally containing one or more heteroatoms selected from sulfur, oxygen, or nitrogen;

R_5' and R_6' are each independently hydrogen or an optionally substituted alkyl;

15 R_3 is hydrogen, halogen, C_{1-4} alkyl, $CH_2NHC(O)C(O)NH_2$, halo-substituted C_{1-4} alkyl, $-CH=CR_8'R_8'$, cyclopropyl optionally substituted by R_8' , CN, OR_8 , CH_2OR_8 , NR_8R_{10} , $CH_2NR_8R_{10}$, $C(Z')H$, $C(O)OR_8$, $C(O)OR_8$, $C(O)NR_8R_{10}$, or $C\equiv CR_8$;

Z' is O, S, NR_8 , NOR_8 , NCN , $C(-CN)_2$, CR_8CN , CR_8NO_2 , $CR_8C(O)OR_8$, $CR_8C(O)ONR_8R_8$, $C(-CN)NO_2$, $C(-CN)C(O)OR_8$, $C(-CN)C(O)NR_8R_8$;

20 Z is $C(Y')R_{14}$, $C(O)OR_{14}$, $C(Y')NR_{10}R_{14}$, $C(NR_{10})NR_{10}R_{14}$, CN, $C(NOR_8)R_{14}$, $C(O)NR_8NR_8C(O)R_8$, $C(O)NR_8NR_{10}R_{14}$, $C(NOR_{14})R_8$, $C(NR_8)NR_{10}R_{14}$, $C(NR_{14})NR_8R_8C(NCN)NR_{10}R_{14}$, $C(NCN)SR_9$, (2-, 4- or 5-imidazolyl), (3-, 4- or 5-pyrazolyl), (4- or 5-triazolyl[1,2,3]), (3- or 5-triazolyl[1,2,4]), (5-tetrazolyl), (2-, 4- or 5-oxazolyl), (3-, 4- or 5-isoxazolyl), (3- or 5-oxadiazolyl[1,2,4]), (2oxadiazolyl[1,3,4]), (2-thiadiazolyl[1,3,4]), (2-, 4-, or 5-thiazolyl), (2-, 4-, or 5-oxazolidinyl), (2-, 4-, or 5-thiazolidinyl), or (2-, 4-, or 5-imidazolidinyl);

25 wherein all of the heterocyclic ring systems may be optionally substituted one or more times by R_{14} ;

the dotted line in the first formula of X_4 represents a single or double bond;

Y' is O or S;

30 R_7 is $-(CR_4R_5)_qR_{12}$ or C_{1-6} alkyl wherein the R_{12} or C_{1-6} alkyl group is optionally substituted one or more times by C_{1-2} alkyl optionally substituted by one to three fluorines, $--F$, $--Br$, $--Cl$, $--NO_2$, $--NR_{10}R_{11}$, $--C(O)R_8$, $--C(O)OR_8$, $--OR_8$, $--CN$, $--C(O)NR_{10}$

R₁₁, --OC(O)NR₁₀ R₁₁, --OC(O)R₈, --NR₁₀ C(O)NR₁₀ R₁₁, --NR₁₀ C(O)R₁₁, --NR₁₀ C(O)OR₉, --NR₁₀ C(O)R₁₃, --C(NR₁₀)NR₁₀ R₁₁, --C(NCN)NR₁₀ R₁₁, --C(NCN)SR₉, --NR₁₀ C(NCN)SR₉, --NR₁₀ C(NCN)NR₁₀ R₁₁, --NR₁₀ S(O)₂ R₉, --S(O)_m R₉, --NR₁₀ C(O)C(O)NR₁₀ R₁₁, --NR₁₀ C(O)C(O)R₁₀, thiazolyl, imidazolyl, oxazolyl, pyrazolyl, triazolyl, or tetrazolyl;

5 q is 0, 1, or 2;

R₁₂ is C₃-C₇-cycloalkyl, (2-, 3- or 4-pyridyl), pyrimidyl, pyrazolyl, (1- or 2-imidazolyl), thiazolyl, triazolyl, pyrrolyl, piperazinyl, piperidiny, morpholinyl, furanyl, (2- or 3-thienyl), (4- or 5-thiazolyl), quinolinyl, naphthyl, or phenyl;

R₈ is independently selected from hydrogen or R₉ ;

10 R₈ ' is R₈ or fluorine;

R₉ is C₁₋₄ alkyl optionally substituted by one to three fluorines;

R₁₀ is OR₈ or R₁₁ ;

R₁₁ is hydrogen, or C₁₋₄ alkyl optionally substituted by one to three fluorines; or when

R₁₀ and R₁₁ are as NR₁₀ R₁₁ they may together with the nitrogen form a 5 to 7 membered

15 ring optionally containing at least one additional heteroatom selected from O/N/or S;

R₁₃ is oxazolidinyl, oxazolyl, thiazolyl, pyrazolyl, triazolyl, tetrazolyl, imidazolyl, imidazolidinyl, thiazolidinyl, isoxazolyl, oxadiazolyl, or thiadiazolyl, and each of these heterocyclic rings is connected through a carbon atom and each may be unsubstituted or substituted by one or two C₁₋₂ alkyl groups;

20 R₁₄ is hydrogen or R₇ ; or when R₁₀ and R₁₄ are as NR₁₀ R₁₄ they may together with the nitrogen form a 5 to 7 membered ring optionally containing one or more additional heteroatoms selected from O, N, or S;

R₁₅ is optionally substituted pyridyl, pyrazinyl, pyrimidinyl, isoxazolyl; or an N-oxide thereof; or a optionally substituted phenyl, wherein s optional substituents are one or

25 more selected from the group consisting of halogen, haloalkyl, aryl, arylalkyl, alkoxy, aryloxy, alkylthio, arylthio, alkoxycarbonyl, alkanoyl, aroyl, alkylsulphonyl, arylsulphonyl, alkylsulphinyl, arylsulphinyl, hydroxy, hydroxyalkyl, formyl, alkanoylamino, aroylamino, cyano, and nitro;

R₁₆ is amino, C₁-C₆ alkylamino, arylamino, C₁-C₆ alkoxy, or aryloxy;

30 R₁₇ is hydrogen, or optionally substituted alkyl, alkenyl, alkoxyalkyl, or alkanoyl, or a formyl, carboxamido, or thiocarboxamido;

R₁₈ is (=O), hydrogen, -C(O)R₂, -CH₃, -CH₂CH₃, -C(O)OH;

provided that:

when R₁₂ is N-pyrazolyl, N-imidazolyl, N-triazolyl, N-pyrrolyl, N-piperazinyl, N-piperidinyl, or N-morpholinyl, then q is not 1; or

5 the pharmaceutically acceptable salts thereof.

31. The method of claim 29, wherein the PDE4 inhibitor is selected from the group consisting of

10 *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof ;

N-(3,5-dichloropyrid-4-yl)-3-cyclopentyloxy-4-methoxybenzamide;

1-[3-cyclopentyloxy)-4-methoxyphenyl] ethanone (E)-O-
{aminocarbonyl}oxime; and

15 (+/-)-4-[2-(3-cyclopentyloxy-4-methoxyphenyl)-2-phenylethyl]pyridine.

32. The method of claim 29, wherein the PDE4 inhibitor is *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.

20 33. The method of claim 29, further comprising administering a therapeutically effective amount of an agent effective in raising intracellular concentrations of cAMP or analogs thereof.

34. The method of claim 33, wherein the agent is salmeterol or salt thereof.

25 35. The method of claim 29, wherein the side effect is myelosuppression.

36. The method of claim 29, wherein the side effect is mucositis.

30 37. The method of claim 29, wherein the side effect is neutropenia

38. Use of (i) an anti-neoplastic agent that has an adverse effect on normal progenitor cells upon use; and (ii) a PDE4 inhibitor in the preparation of a medicament for the treatment of cancer.

5 39. The use of claim 38, wherein the anti-neoplastic agent is a phase specific anti-neoplastic agent.

40. The use of claim 39, wherein the phase specific anti-neoplastic agent is selected from the group consisting of diterpenoids; vinca alkaloids; epipodo -phyllotoxins,
10 antimetabolites, camptothecins, or pharmaceutically acceptable salts thereof.

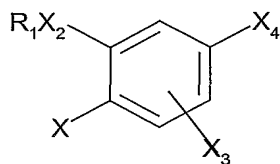
41. The use of claim 38, wherein the anti-neoplastic agent is paclitaxel.

42. The use of claim 38, wherein the anti-neoplastic agent is topotecan.
15

43. The use of claim 38, wherein the PDE4 inhibitor is a compound of formula I

20

I



wherein: R_1 is $-(CR_4 R_5)_n C(O)O(CR_4 R_5)_m R_6$, $-(CR_4 R_5)_n C(O)NR_4 (CR_4 R_5)_m R_6$, $-(CR_4 R_5)_n$
25 $O(CR_4 R_5)_m R_6$, or $--(CR_4 R_5)_r R_6$ wherein the alkyl moieties may be optionally substituted
with one or more halogens;
 m is 0 to 2;

n is 0 to 4;

r is 0 to 6;

R₄ and R₅ are independently selected from hydrogen or a C₁₋₂ alkyl;

R₆ is hydrogen, methyl, hydroxyl, aryl, halo substituted aryl, aryloxy C₁₋₃ alkyl, halo

- 5 substituted aryloxy C₁₋₃ alkyl, indanyl, indenyl, C₇₋₁₁ polycycloalkyl, tetrahydrofuranyl, furanyl, tetrahydropyranyl, pyranyl, tetrahydrothienyl, thienyl, tetrahydrothiopyranyl, thiopyranyl, C₃₋₆ cycloalkyl, or a C₄₋₆ cycloalkyl containing one or two unsaturated bonds, wherein the cycloalkyl and heterocyclic moieties may be optionally substituted by 1 to 3 methyl groups or one ethyl group;

- 10 provided that:

when R₆ is hydroxyl, then m is 2; or

when R₆ is hydroxyl, then r is 2 to 6; or

when R₆ is 2-tetrahydropyranyl, 2-tetrahydrothiopyranyl, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then m is 1 or 2; or

- 15 when R₆ is 2-tetrahydropyranyl, 2-tetrahydrothiopyranyl, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then r is 1 to 6;

when n is 1 and m is 0, then R₆ is other than H in $--(CR_4 R_5)_n O(CR_4 R_5)_m R_6$

;

X is YR₂, halogen, nitro, NR₄ R₅, or formyl amine;

- 20 Y is O or S(O)_{m'};

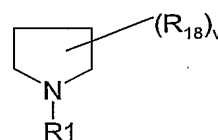
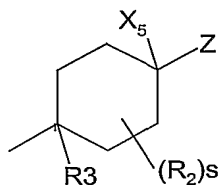
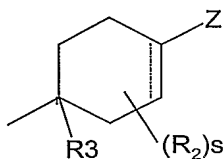
m' is 0, 1, or 2;

X₂ is O or NR₈;

X₃ is hydrogen or X;

X₄ is

25



$C(Z')NHR_{15}$, $CR_4(NOC(O)R_{16})$, or $CR_2'R_3'CR_4'R_5'R_6'$;

- 5 X_5 is H, R_9 , OR_8 , CN, $C(O)R_8$, $C(O)OR_8$, $C(O)NR_8R_8$, or NR_8R_8 ;
 R_2 is independently selected from the group consisting of $--CH_3$ and $--CH_2CH_3$
optionally substituted by 1 or more halogens;
 R_2' is a hydrogen, halogen, or OR_{17} ;
s is 0 to 4;
- 10 R_3' and R_4' are each independently $-(CH_2)_tX_6$;
t is 0, 1, 2, or 3;
v is 0, 1, 2, or 3;
 X_6 is a mono- or bicyclic aryl group optionally containing one or more heteroatoms
selected from sulfur, oxygen, or nitrogen;
- 15 R_5' and R_6' are each independently hydrogen or an optionally substituted alkyl;
 R_3 is hydrogen, halogen, C_{1-4} alkyl, $CH_2NHC(O)C(O)NH_2$, halo-substituted C_{1-4} alkyl, $-CH=CR_8'R_8'$, cyclopropyl optionally substituted by R_8' , CN, OR_8 , CH_2OR_8 , NR_8R_{10} ,
 $CH_2NR_8R_{10}$, $C(Z')H$, $C(O)OR_8$, $C(O)OR_8$, $C(O)NR_8R_{10}$, or $C\equiv CR_8$;
 Z' is O, S, NR_8 , NOR_8 , NCN , $C(-CN)_2$, CR_8CN , CR_8NO_2 , $CR_8C(O)OR_8$, $CR_8C(O)ONR_8R_8$, $C(-$
20 $CN)NO_2$, $C(-CN)C(O)OR_9$, $C(-CN)C(O)NR_8R_8$;
Z is $C(Y')R_{14}$, $C(O)OR_{14}$, $C(Y')NR_{10}R_{14}$, $C(NR_{10})NR_{10}R_{14}$, CN, $C(NOR_8)R_{14}$, $C(O)NR_8NR_8C(O)R_8$,
 $C(O)NR_8NR_{10}R_{14}$, $C(NOR_{14})R_8$, $C(NR_8)NR_{10}R_{14}$, $C(NR_{14})NR_8R_8C(NCN)NR_{10}R_{14}$, $C(NCN)SR_9$,
(2-, 4- or 5-imidazolyl), (3-, 4- or 5-pyrazolyl), (4- or 5-triazolyl[1,2,3]), (3- or 5-
25 triazolyl[1,2,4]), (5-tetrazolyl), (2-, 4- or 5-oxazolyl), (3-, 4- or 5-isoxazolyl), (3- or 5-
oxadiazolyl[1,2,4]), (2oxadiazolyl[1,3,4]), (2-thiadiazolyl[1,3,4]), (2-, 4-, or 5-thiazolyl),
(2-, 4-, or 5-oxazolidinyl), (2-, 4-, or 5-thiazolidinyl), or (2-, 4-, or 5-imidazolidinyl);
wherein all of the heterocyclic ring systems may be optionally substituted one or more
times by R_{14} ;
the dotted line in the first formula of X_4 represents a single or double bond;
- 30 Y' is O or S;

- R₇ is --(CR₄ R₅)_q R₁₂ or C₁₋₆ alkyl wherein the R₁₂ or C₁₋₆ alkyl group is optionally substituted one or more times by C₁₋₂ alkyl optionally substituted by one to three fluorines, --F, --Br, --Cl, --NO₂, --NR₁₀ R₁₁, --C(O)R₈, --C(O)OR₈, --OR₈, --CN, --C(O)NR₁₀ R₁₁, --OC(O)NR₁₀ R₁₁, --OC(O)R₈, --NR₁₀ C(O)NR₁₀ R₁₁, --NR₁₀ C(O)R₁₁, --NR₁₀ C(O)OR₉, --NR₁₀ C(O)R₁₃, --C(NR₁₀)NR₁₀ R₁₁, --C(NCN)NR₁₀ R₁₁, --C(NCN)SR₉, --NR₁₀ C(NCN)SR₉, --NR₁₀ C(NCN)NR₁₀ R₁₁, --NR₁₀ S(O)₂ R₉, --S(O)_m R₉, --NR₁₀ C(O)C(O)NR₁₀ R₁₁, --NR₁₀ C(O)C(O)R₁₀, thiazolyl, imidazolyl, oxazolyl, pyrazolyl, triazolyl, or tetrazolyl; q is 0, 1, or 2;
- R₁₂ is C₃ -C₇ -cycloalkyl, (2-, 3- or 4-pyridyl), pyrimidyl, pyrazolyl, (1- or 2-imidazolyl), thiazolyl, triazolyl, pyrrolyl, piperazinyl, piperidinyl, morpholinyl, furanyl, (2- or 3-thienyl), (4- or 5-thiazolyl), quinolinyl, naphthyl, or phenyl;
- R₈ is independently selected from hydrogen or R₉ ;
- R₈ ' is R₈ or fluorine;
- R₉ is C₁₋₄ alkyl optionally substituted by one to three fluorines;
- R₁₀ is OR₈ or R₁₁ ;
- R₁₁ is hydrogen, or C₁₋₄ alkyl optionally substituted by one to three fluorines; or when R₁₀ and R₁₁ are as NR₁₀ R₁₁ they may together with the nitrogen form a 5 to 7 membered ring optionally containing at least one additional heteroatom selected from O/N/or S;
- R₁₃ is oxazolidinyl, oxazolyl, thiazolyl, pyrazolyl, triazolyl, tetrazolyl, imidazolyl, imidazolidinyl, thiazolidinyl, isoxazolyl, oxadiazolyl, or thiadiazolyl, and each of these heterocyclic rings is connected through a carbon atom and each may be unsubstituted or substituted by one or two C₁₋₂ alkyl groups;
- R₁₄ is hydrogen or R₇ ; or when R₁₀ and R₁₄ are as NR₁₀ R₁₄ they may together with the nitrogen form a 5 to 7 membered ring optionally containing one or more additional heteroatoms selected from O, N, or S;
- R₁₅ is optionally substituted pyridyl, pyrazinyl, pyrimidinyl, isoxazolyl; or an N-oxide thereof; or a optionally substituted phenyl, wherein s optional substituents are one or more selected from the group consisting of halogen, haloalkyl, aryl, arylalkyl, alkoxy, aryloxy, alkylthio, arylthio, alkoxycarbonyl, alkanoyl, aroyl, alkylsulphonyl, arylsulphonyl, alkylsulphinyl, arylsulphinyl, hydroxy, hydroxyalkyl, formyl, alkanoylamino, aroylamino, cyano, and nitro;

R₁₆ is amino, C₁-C₆ alkylamino, arylamino, C₁-C₆ alkoxy, or aryloxy;

R₁₇ is hydrogen, or optionally substituted alkyl, alkenyl, alkoxyalkyl, or alkanoyl, or a formyl, carboxamido, or thiocarboxamido;

R₁₈ is (=O), hydrogen, -C(O)R₂, -CH₃, -CH₂CH₃, -C(O)OH;

5 provided that:

when R₁₂ is N-pyrazolyl, N-imidazolyl, N-triazolyl, N-pyrrolyl, N-piperazinyl, N-piperidinyl, or N-morpholinyl, then q is not 1; or
the pharmaceutically acceptable salts thereof.

10 44. The use of claim 38, wherein the PDE4 inhibitor is selected from the group consisting of
cis-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof;

N-(3,5-dichloropyrid-4-yl)-3-cyclopentyloxy-4-methoxybenzamide;
15 1-[3-cyclopentyloxy)-4-methoxyphenyl] ethanone (E)-O-
{aminocarbonyl} oxime; and
(+/-)-4-[2-(3-cyclopentyloxy-4-methoxyphenyl)-2-phenylethyl]pyridine.

45. The use of claim 38, wherein the PDE4 inhibitor is *cis*-4-cyano-4-[3-
20 (cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.

46. The use of claim 38, further comprising administering a therapeutically effective amount of an agent effective in raising intracellular concentrations of cAMP or analogs thereof.

25

47. The use of claim 46, wherein the agent is salmeterol or a salt thereof.

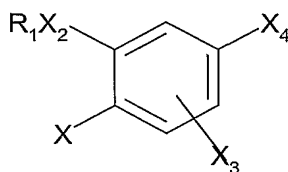
48. Use of a PDE4 inhibitor for preparing a medicament for attenuating cancer treatment side effects on normal progenitor cells.

30

49. The use of claim 48, wherein the PDE4 inhibitor is a compound of formula I

5

I



wherein: R₁ is $-(\text{CR}_4 \text{ R}_5)_n \text{C}(\text{O})\text{O}(\text{CR}_4 \text{ R}_5)_m \text{R}_6$, $-(\text{CR}_4 \text{ R}_5)_n \text{C}(\text{O})\text{NR}_4 (\text{CR}_4 \text{ R}_5)_m \text{R}_6$, $-(\text{CR}_4 \text{ R}_5)_n \text{O}(\text{CR}_4 \text{ R}_5)_m \text{R}_6$, or $--(\text{CR}_4 \text{ R}_5)_r \text{R}_6$ wherein the alkyl moieties may be optionally substituted with one or more halogens;

10

m is 0 to 2;

n is 0 to 4;

r is 0 to 6;

R₄ and R₅ are independently selected from hydrogen or a C₁₋₂ alkyl;

15

R₆ is hydrogen, methyl, hydroxyl, aryl, halo substituted aryl, aryloxy C₁₋₃ alkyl, halo substituted aryloxy C₁₋₃ alkyl, indanyl, indenyl, C₇₋₁₁ polycycloalkyl, tetrahydrofuranyl, furanyl, tetrahydropyranyl, pyranal, tetrahydrothienyl, thienyl, tetrahydrothiopyranal, thiopyranal, C₃₋₆ cycloalkyl, or a C₄₋₆ cycloalkyl containing one or two unsaturated bonds, wherein the cycloalkyl and heterocyclic moieties may be optionally substituted by 1 to 3 methyl groups or one ethyl group;

20

provided that:

when R₆ is hydroxyl, then m is 2; or

when R₆ is hydroxyl, then r is 2 to 6; or

when R₆ is 2-tetrahydropyranyl, 2-tetrahydrothiopyranal, 2-

25

tetrahydrofuranyl, or 2-tetrahydrothienyl, then m is 1 or 2; or

when R₆ is 2-tetrahydropyranyl, 2-tetrahydrothiopyranal, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then r is 1 to 6;

when n is 1 and m is 0, then R₆ is other than H in $--(CR_4 R_5)_n O(CR_4 R_5)_m R_6$

;

X is YR₂, halogen, nitro, NR₄ R₅, or formyl amine;

Y is O or S(O)_{m'};

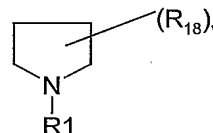
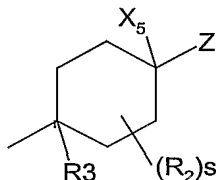
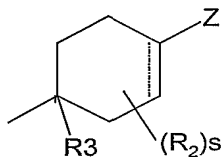
5 m' is 0, 1, or 2;

X₂ is O or NR₈;

X₃ is hydrogen or X;

X₄ is

10



15 C(Z')NHR₁₅, CR₄(NOC(O)R₁₆), or CR₂'R₃'CR₄'R₅'R₆';

X₅ is H, R₉, OR₈, CN, C(O)R₈, C(O)OR₈, C(O)NR₈ R₈, or NR₈ R₈;

R₂ is independently selected from the group consisting of $--CH_3$ and $--CH_2 CH_3$ optionally substituted by 1 or more halogens;

20 R₂' is a hydrogen, halogen, or OR₁₇;

s is 0 to 4;

R₃' and R₄' are each independently $-(CH_2)_t X_6$;

t is 0, 1, 2, or 3;

v is 0, 1, 2, or 3;

25 X₆ is a mono- or bicyclic aryl group optionally containing one or more heteroatoms selected from sulfur, oxygen, or nitrogen;

- R_5' and R_6' are each independently hydrogen or an optionally substituted alkyl;
 R_3 is hydrogen, halogen, C_{1-4} alkyl, $CH_2NHC(O)C(O)NH_2$, halo-substituted C_{1-4} alkyl, $-CH=CR_8'R_8'$, cyclopropyl optionally substituted by R_8' , CN, OR_8 , CH_2OR_8 , NR_8R_{10} , $CH_2NR_8R_{10}$, $C(Z')H$, $C(O)OR_8$, $C(O)OR_8$, $C(O)NR_8R_{10}$, or $C\equiv CR_8$;
- 5 Z' is O, S, NR_8 , NOR_8 , NCN , $C(-CN)_2$, CR_8CN , CR_8NO_2 , $CR_8C(O)OR_8$, $CR_8C(O)ONR_8R_8$, $C(-CN)NO_2$, $C(-CN)C(O)OR_9$, $C(-CN)C(O)NR_8R_8$;
 Z is $C(Y')R_{14}$, $C(O)OR_{14}$, $C(Y')NR_{10}R_{14}$, $C(NR_{10})NR_{10}R_{14}$, CN, $C(NOR_8)R_{14}$, $C(O)NR_8NR_8C(O)R_8$, $C(O)NR_8NR_{10}R_{14}$, $C(NOR_{14})R_8$, $C(NR_8)NR_{10}R_{14}$, $C(NR_{14})NR_8R_8C(NCN)NR_{10}R_{14}$, $C(NCN)SR_9$, (2-, 4- or 5-imidazolyl), (3-, 4- or 5-pyrazolyl), (4- or 5-triazolyl[1,2,3]), (3- or 5-
 10 triazolyl[1,2,4]), (5-tetrazolyl), (2-, 4- or 5-oxazolyl), (3-, 4- or 5-isoxazolyl), (3- or 5-oxadiazolyl[1,2,4]), (2oxadiazolyl[1,3,4]), (2-thiadiazolyl[1,3,4]), (2-, 4-, or 5-thiazolyl), (2-, 4-, or 5-oxazolidinyl), (2-, 4-, or 5-thiazolidinyl), or (2-, 4-, or 5-imidazolidinyl);
 wherein all of the heterocyclic ring systems may be optionally substituted one or more times by R_{14} ;
- 15 the dotted line in the first formula of X_4 represents a single or double bond;
 Y' is O or S;
 R_7 is $-(CR_4R_5)_qR_{12}$ or C_{1-6} alkyl wherein the R_{12} or C_{1-6} alkyl group is optionally substituted one or more times by C_{1-2} alkyl optionally substituted by one to three fluorines, $--F$, $--Br$, $--Cl$, $--NO_2$, $--NR_{10}R_{11}$, $--C(O)R_8$, $--C(O)OR_8$, $--OR_8$, $--CN$, $--C(O)NR_{10}$
 20 R_{11} , $--OC(O)NR_{10}R_{11}$, $--OC(O)R_8$, $--NR_{10}C(O)NR_{10}R_{11}$, $--NR_{10}C(O)R_{11}$, $--NR_{10}C(O)OR_9$, $--NR_{10}C(O)R_{13}$, $--C(NR_{10})NR_{10}R_{11}$, $--C(NCN)NR_{10}R_{11}$, $--C(NCN)SR_9$, $--NR_{10}C(NCN)SR_9$, $--NR_{10}C(NCN)NR_{10}R_{11}$, $--NR_{10}S(O)_2R_9$, $--S(O)_mR_9$, $--NR_{10}C(O)C(O)NR_{10}R_{11}$, $--NR_{10}C(O)C(O)R_{10}$, thiazolyl, imidazolyl, oxazolyl, pyrazolyl, triazolyl, or tetrazolyl;
 q is 0, 1, or 2;
- 25 R_{12} is C_3-C_7 -cycloalkyl, (2-, 3- or 4-pyridyl), pyrimidyl, pyrazolyl, (1- or 2-imidazolyl), thiazolyl, triazolyl, pyrrolyl, piperaziny, piperidiny, morpholiny, furanyl, (2- or 3-thienyl), (4- or 5-thiazolyl), quinoliny, naphthyl, or phenyl;
 R_8 is independently selected from hydrogen or R_9 ;
- R_8' is R_8 or fluorine;
- 30 R_9 is C_{1-4} alkyl optionally substituted by one to three fluorines;
 R_{10} is OR_8 or R_{11} ;

R₁₁ is hydrogen, or C₁₋₄ alkyl optionally substituted by one to three fluorines; or when R₁₀ and R₁₁ are as NR₁₀ R₁₁ they may together with the nitrogen form a 5 to 7 membered ring optionally containing at least one additional heteroatom selected from O/N/or S;

R₁₃ is oxazolidinyl, oxazolyl, thiazolyl, pyrazolyl, triazolyl, tetrazolyl, imidazolyl,

5 imidazolidinyl, thiazolidinyl, isoxazolyl, oxadiazolyl, or thiadiazolyl, and each of these heterocyclic rings is connected through a carbon atom and each may be unsubstituted or substituted by one or two C₁₋₂ alkyl groups;

R₁₄ is hydrogen or R₇ ; or when R₁₀ and R₁₄ are as NR₁₀ R₁₄ they may together with the nitrogen form a 5 to 7 membered ring optionally containing one or more additional

10 heteroatoms selected from O, N, or S;

R₁₅ is optionally substituted pyridyl, pyrazinyl, pyrimidinyl, isoxazolyl; or an N-oxide thereof; or a optionally substituted phenyl, wherein s optional substituents are one or more selected from the group consisting of halogen, haloalkyl, aryl, arylalkyl, alkoxy, aryloxy, alkylthio, arylthio, alkoxycarbonyl, alkanoyl, aroyl, alkylsulphonyl, arylsulphonyl, 15 alkylsulphinyl, arylsulphinyl, hydroxy, hydroxyalkyl, formyl, alkanoylamino, aroylamino, cyano, and nitro;

R₁₆ is amino, C₁-C₆ alkylamino, arylamino, C₁-C₆ alkoxy, or aryloxy;

R₁₇ is hydrogen, or optionally substituted alkyl, alkenyl, alkoxyalkyl, or alkanoyl, or a formyl, carboxamido, or thiocarboxamido;

20 R₁₈ is (=O), hydrogen, -C(O)R₂, -CH₃, -CH₂CH₃, -C(O)OH; provided that:

when R₁₂ is N-pyrazolyl, N-imidazolyl, N-triazolyl, N-pyrrolyl, N-piperazinyl, N-piperidinyl, or N-morpholinyl, then q is not 1; or the pharmaceutically acceptable salts thereof.

25

50. The use of claim 48, wherein the PDE4 inhibitor is selected from the group consisting of

cis-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof;

30 N-(3,5-dichloropyrid-4-yl)-3-cyclopentyloxy-4-methoxybenzamide;

1-[3-cyclopentyloxy)-4-methoxyphenyl] ethanone (E)-O-
{aminocarbonyl}oxime; and
(+/-)-4-[2-(3-cyclopentyloxy-4-methoxyphenyl)-2-phenylethyl]pyridine.

- 5 51. The use of claim 48, wherein the PDE4 inhibitor is *cis*-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexane-1-carboxylic acid or salt thereof.
52. The use of claim 48, further comprising administering a therapeutically effective amount of an agent effective in raising intracellular concentrations of cAMP or analogs
10 thereof.
53. The use of claim 48, wherein the agent is salmeterol or salt thereof.
54. A method of treating cancer in a mammal, comprising: administering to the
15 mammal a therapeutically effective amount of a PDE4 inhibitor.

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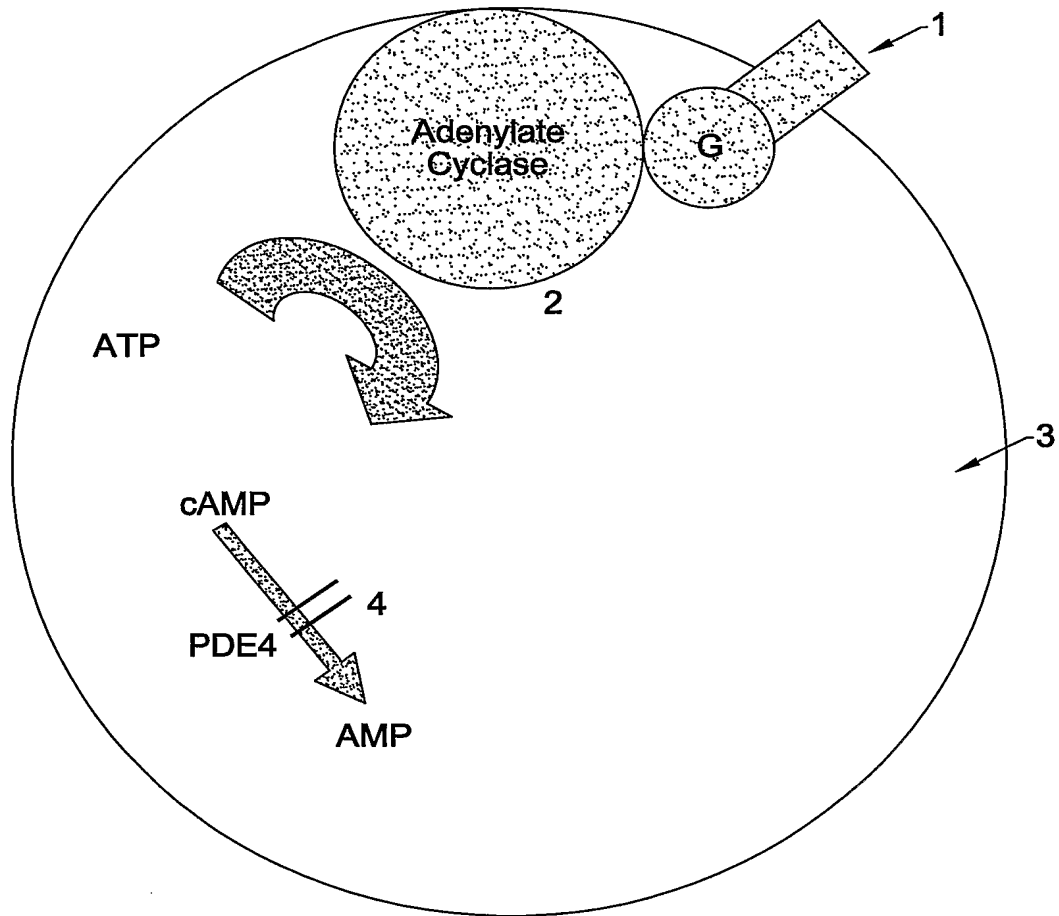
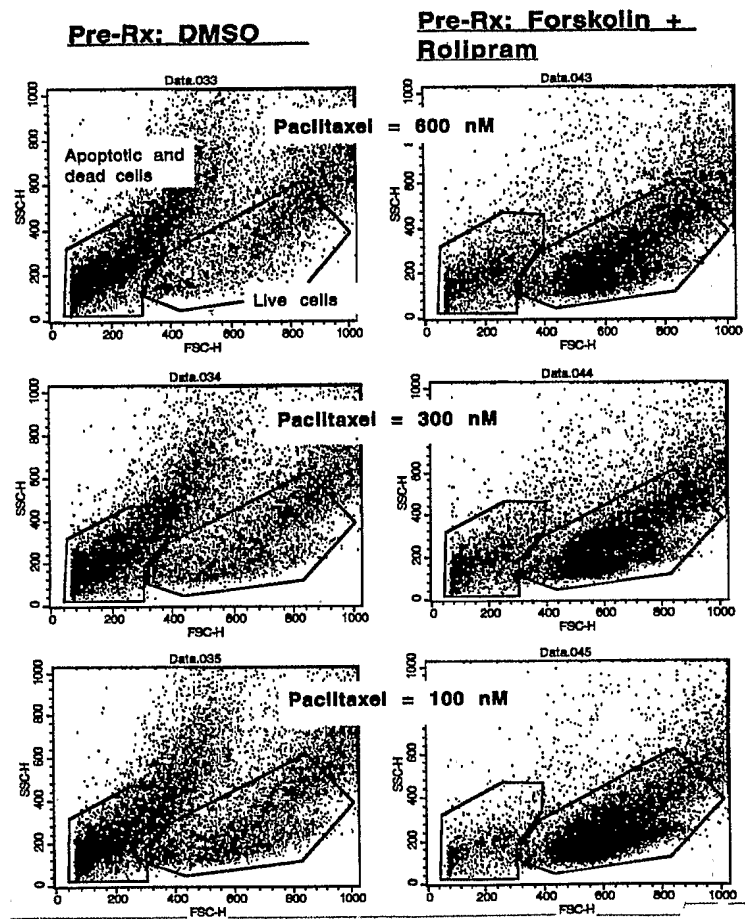
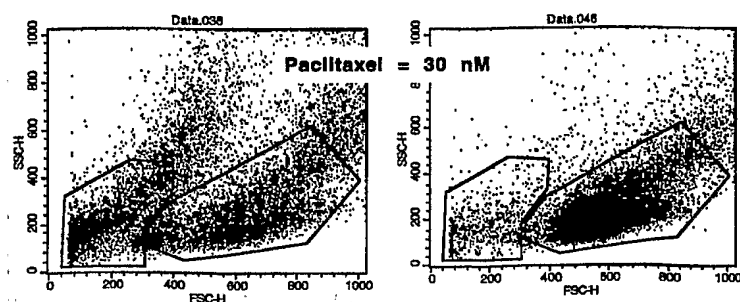
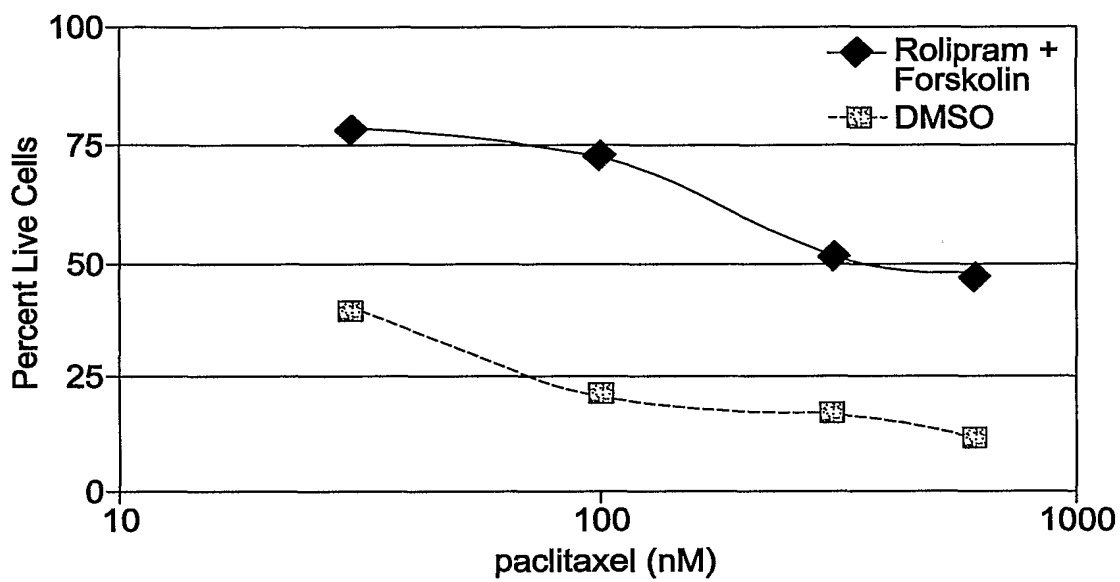


FIG. 1.

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**FIG. 2(a).****FIG. 2(b).**

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FIG. 2(c).

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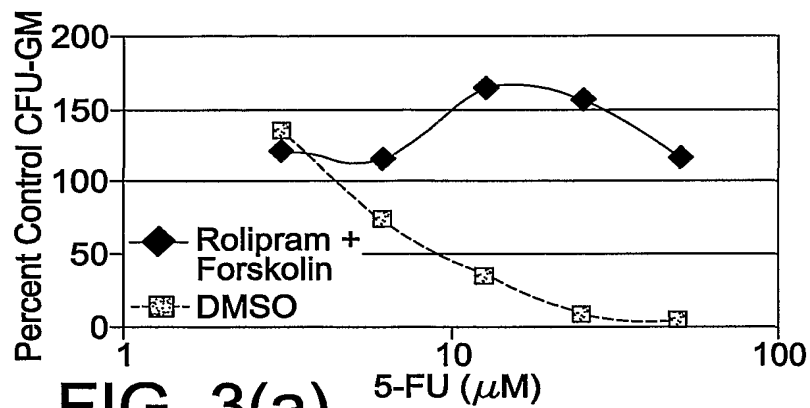


FIG. 3(a).

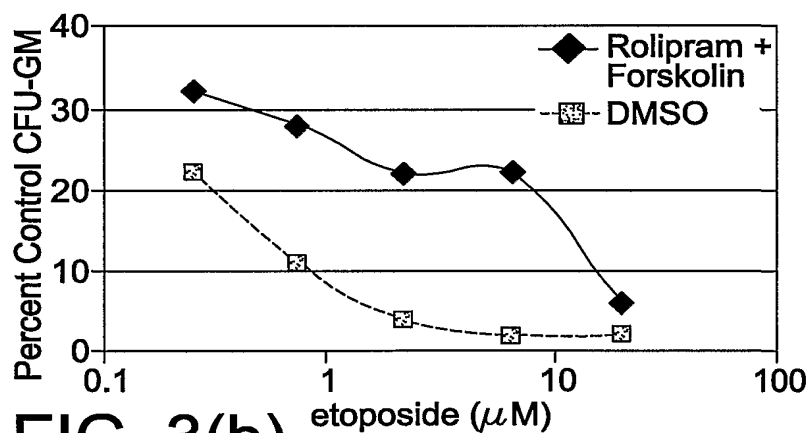


FIG. 3(b).

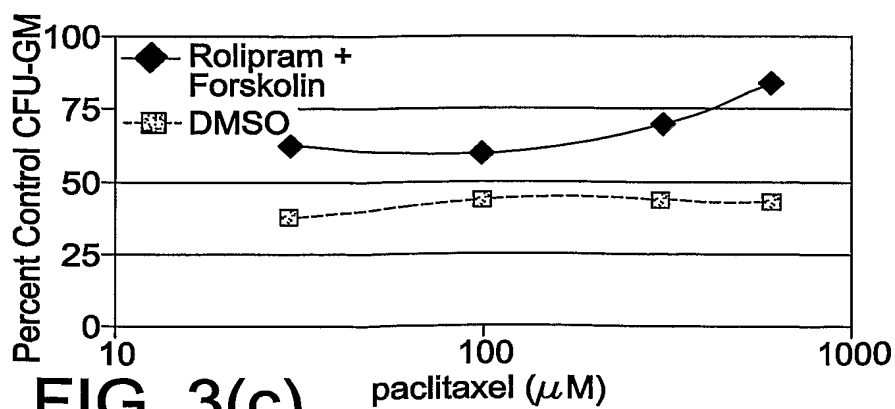
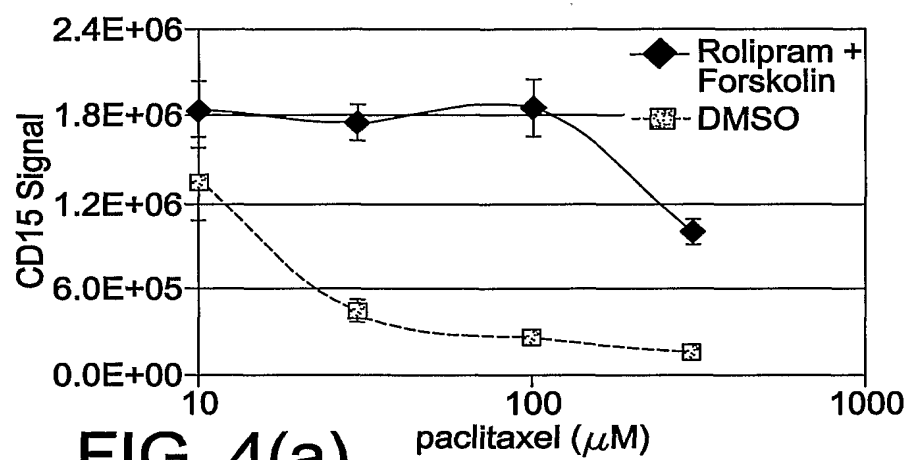
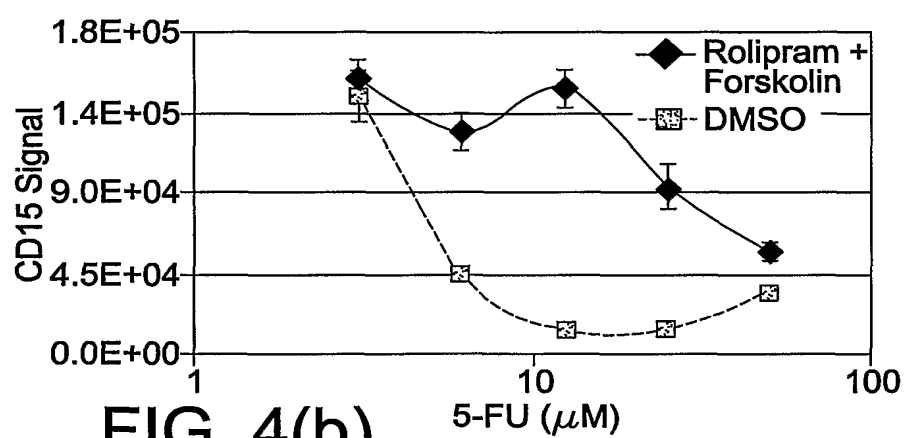
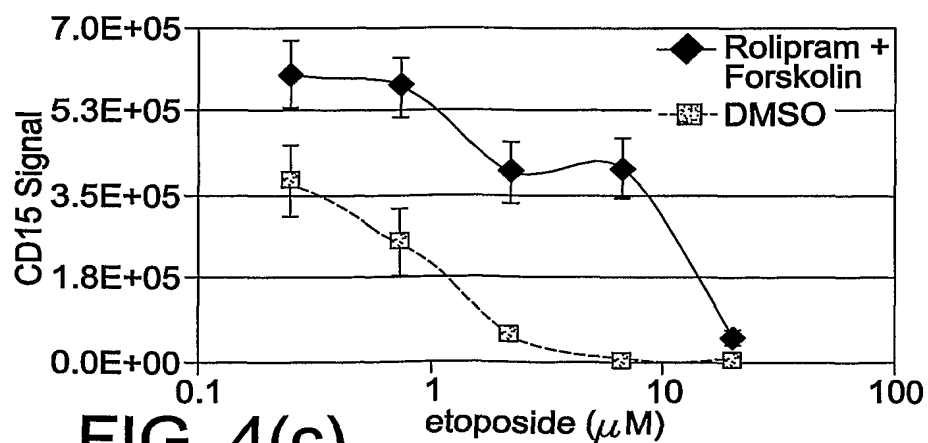
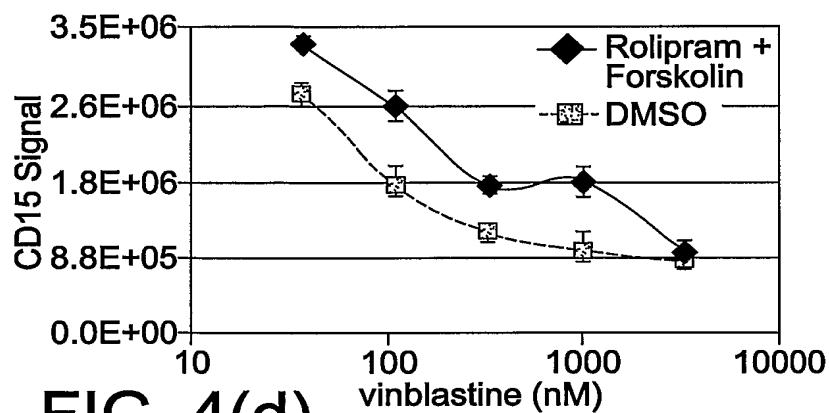
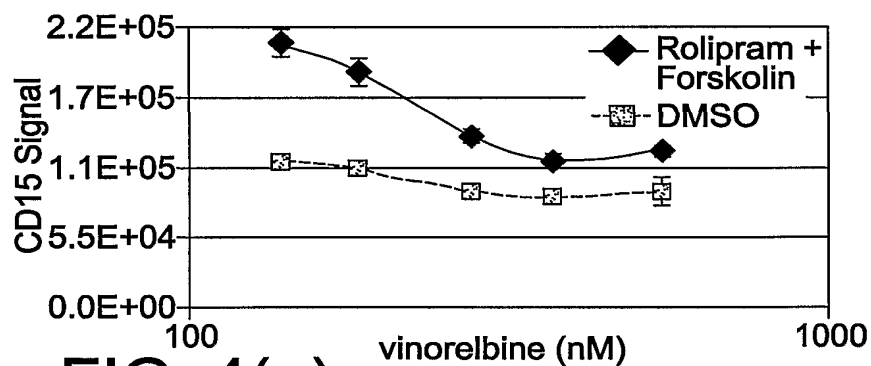
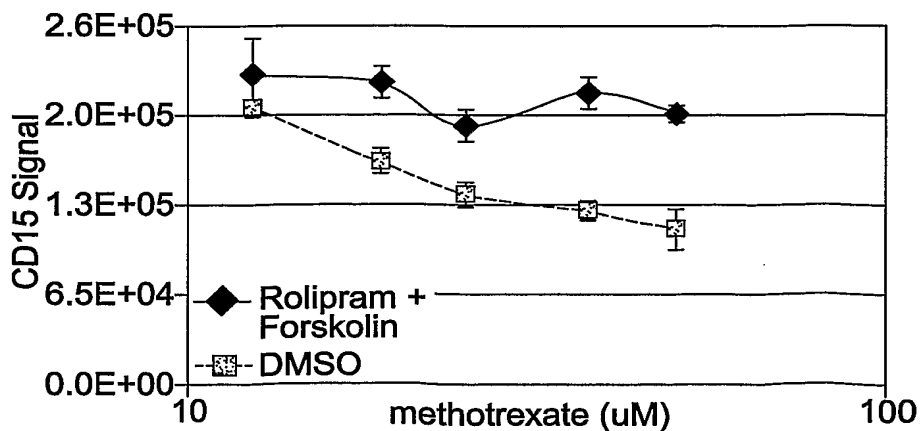


FIG. 3(c).

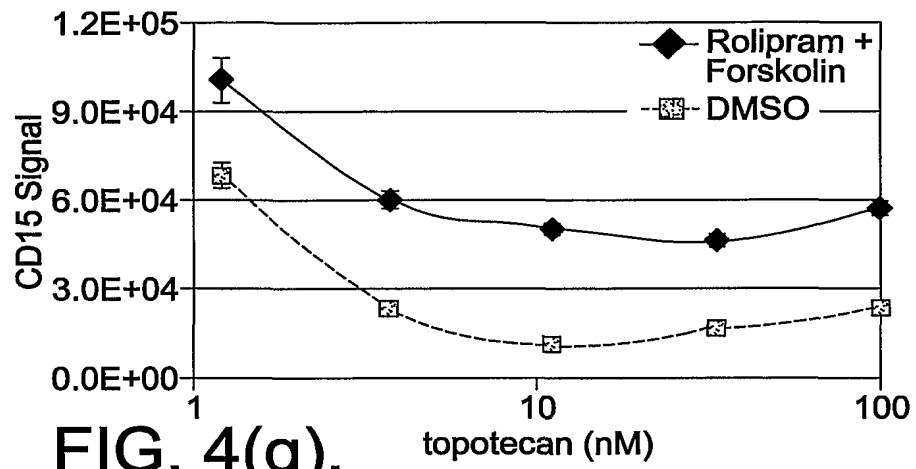
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FIG. 4(a).FIG. 4(b).FIG. 4(c).

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FIG. 4(d).FIG. 4(e).FIG. 4(f).

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**FIG. 4(g).**

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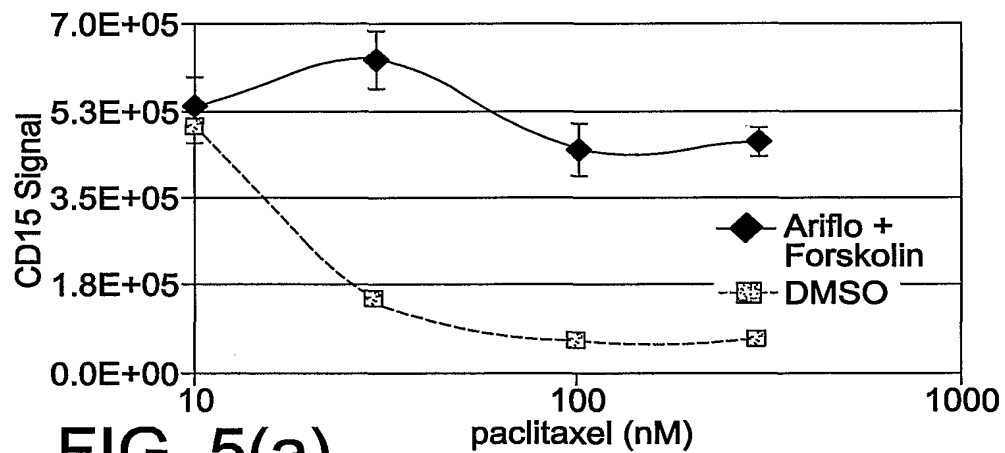


FIG. 5(a).

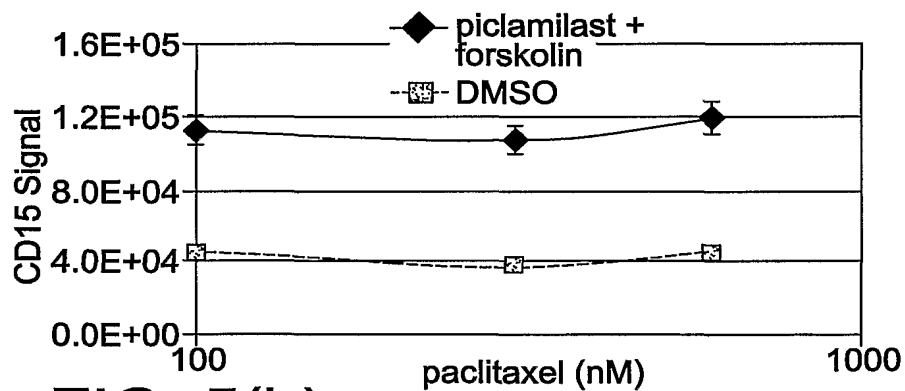


FIG. 5(b).

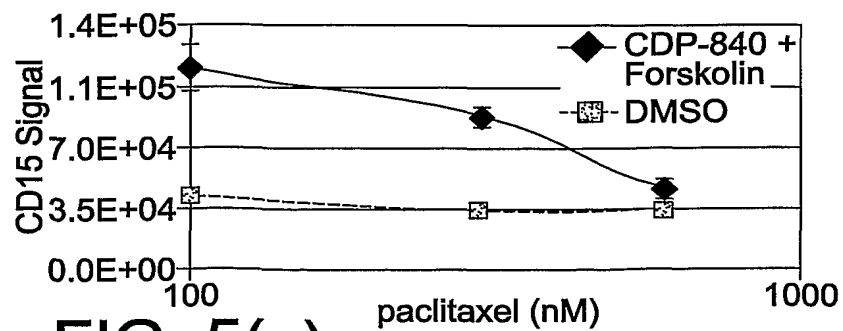


FIG. 5(c).

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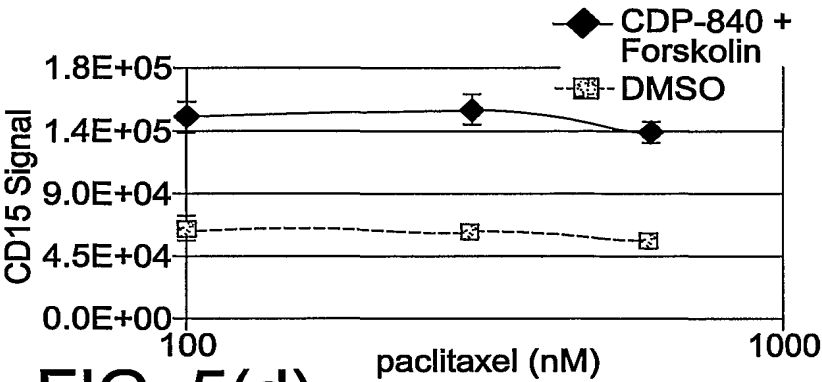
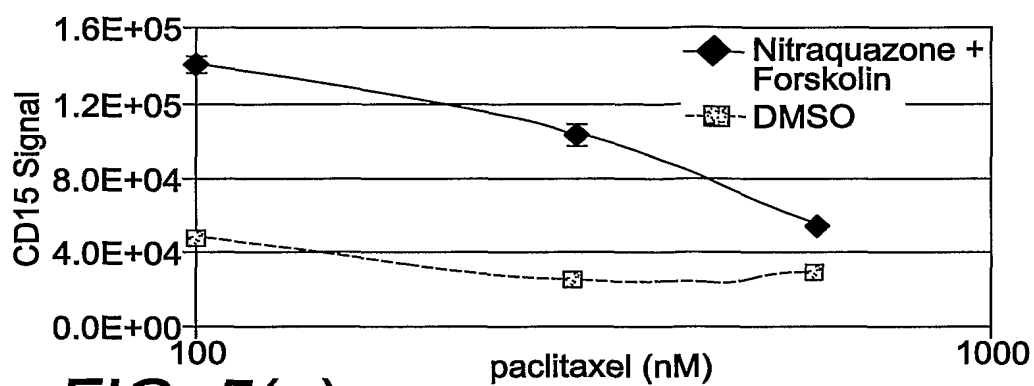
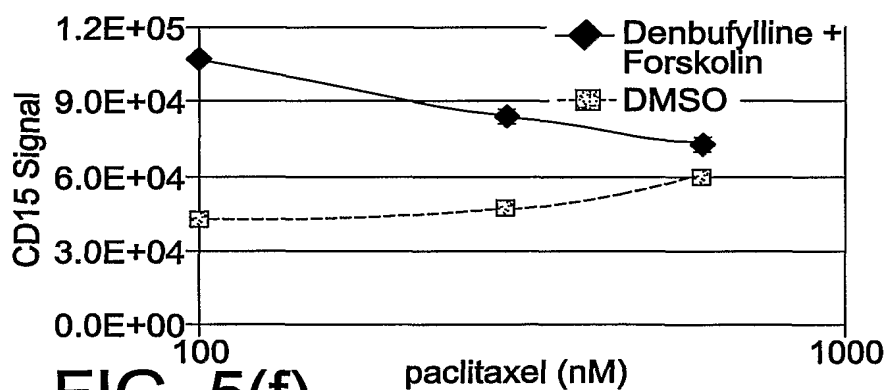
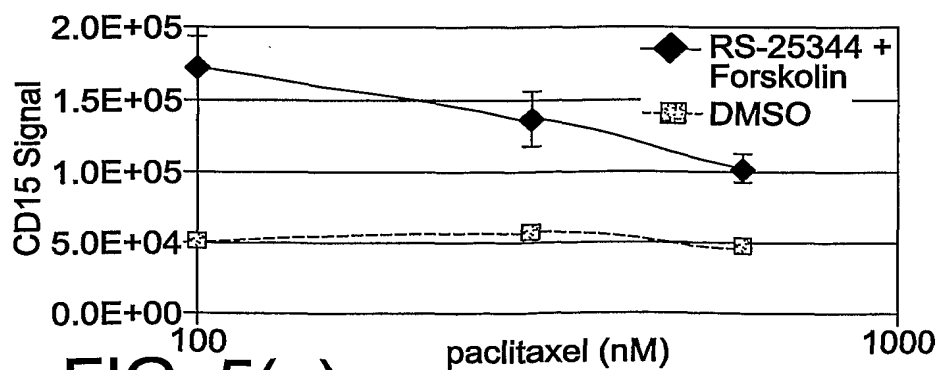
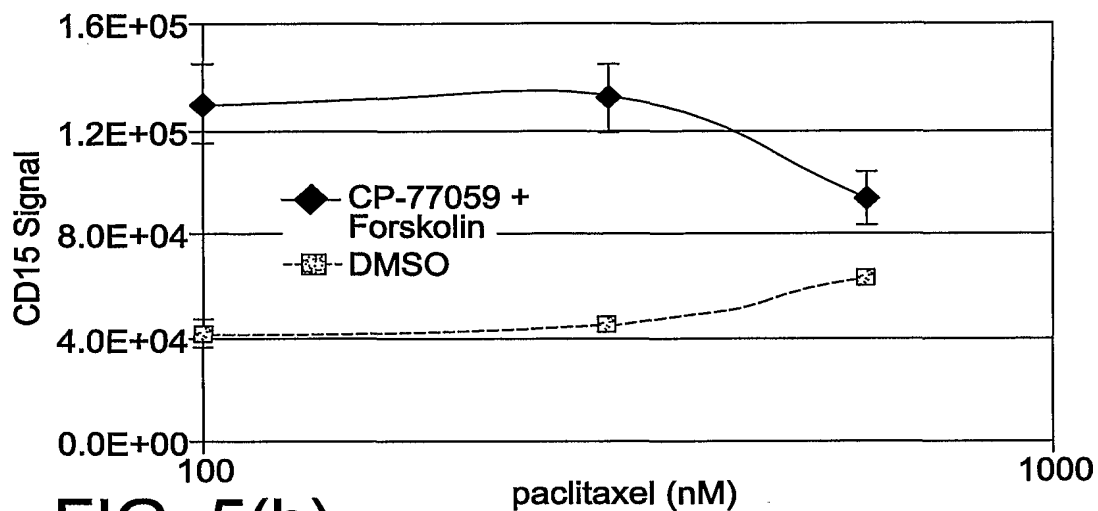
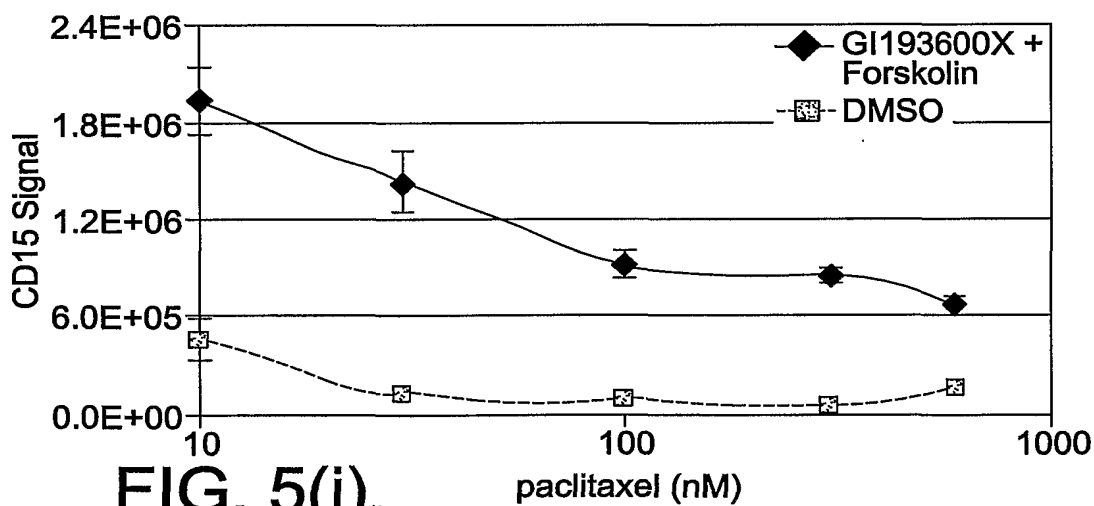
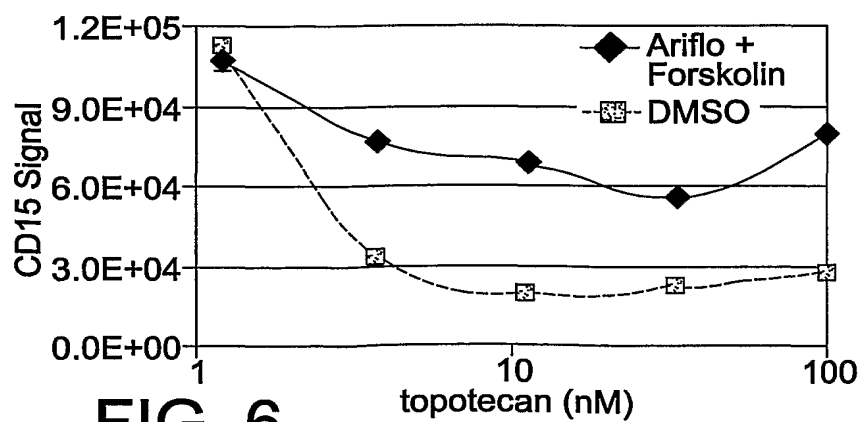


FIG. 5(d).

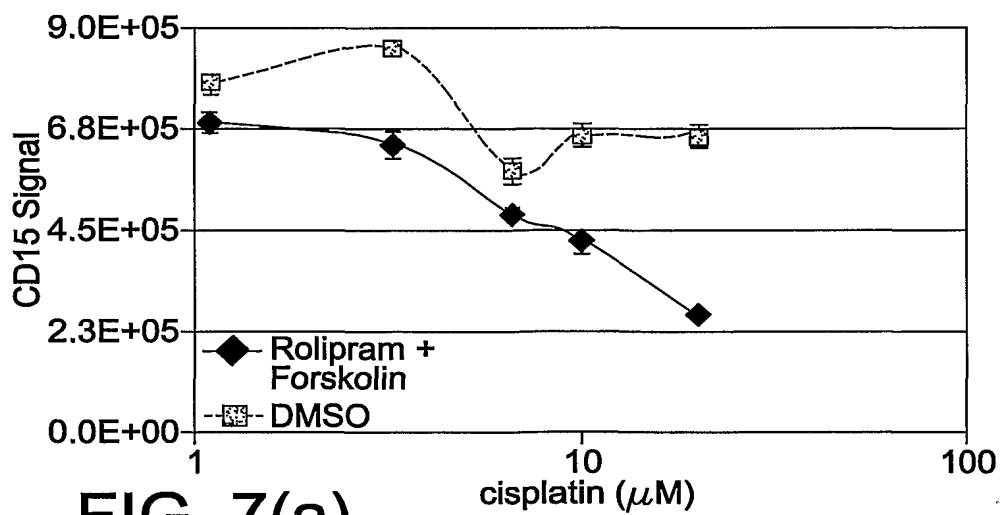
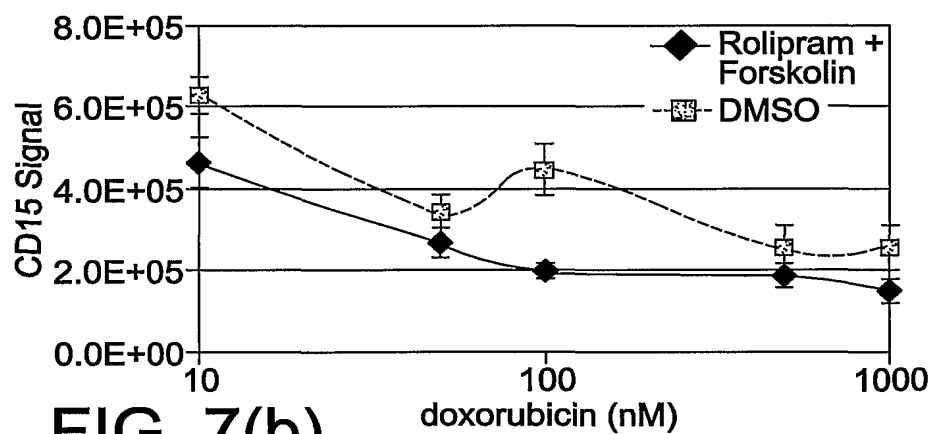
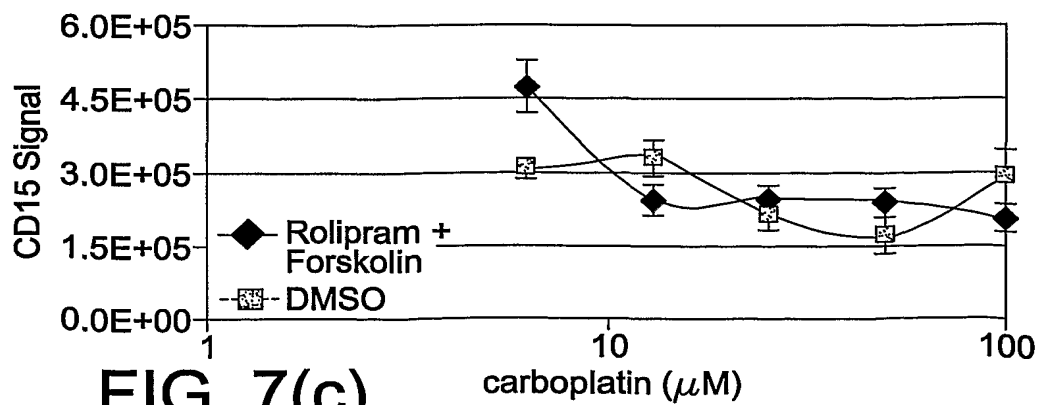
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FIG. 5(e).FIG. 5(f).FIG. 5(g).

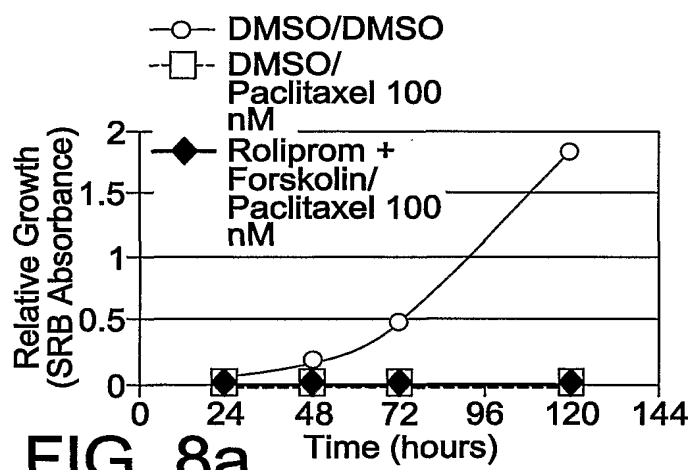
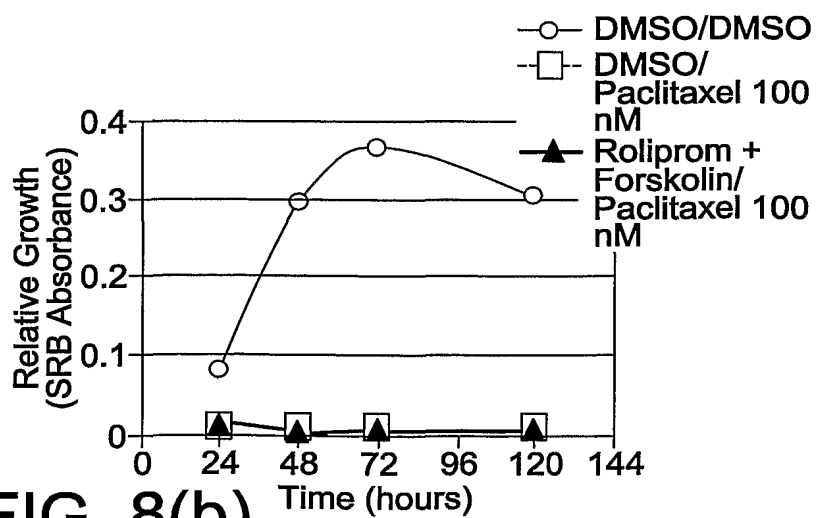
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**FIG. 5(h).****FIG. 5(i).****FIG. 6.**

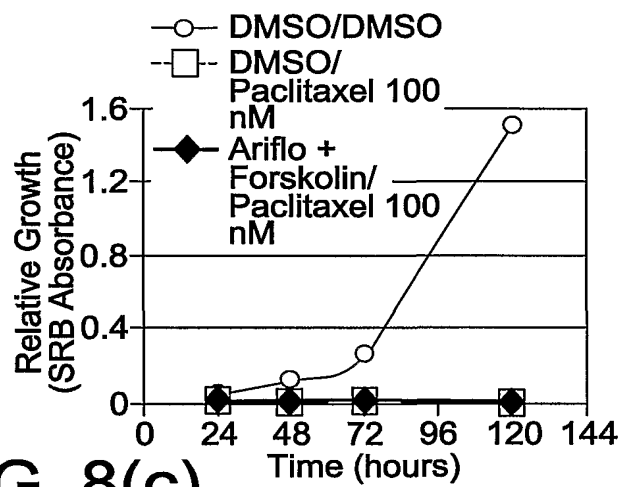
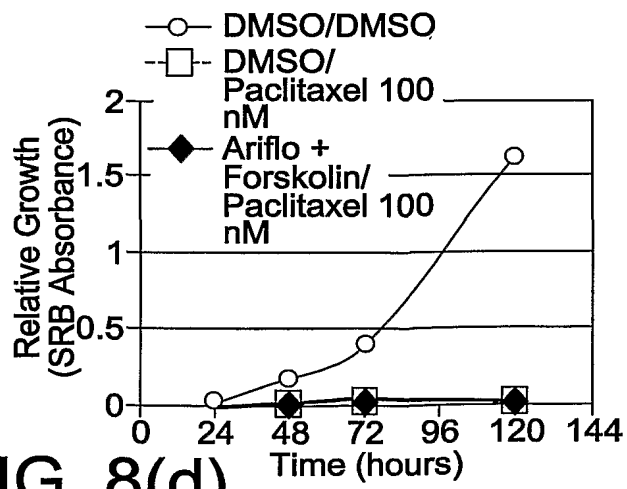
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**FIG. 7(a).****FIG. 7(b).****FIG. 7(c).**

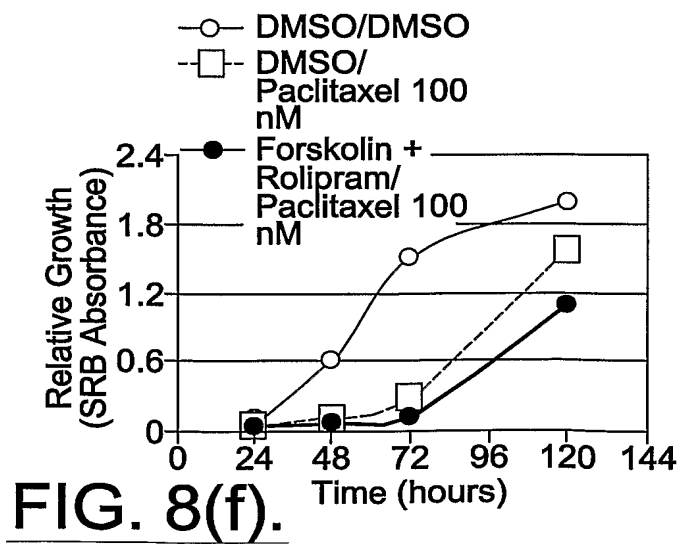
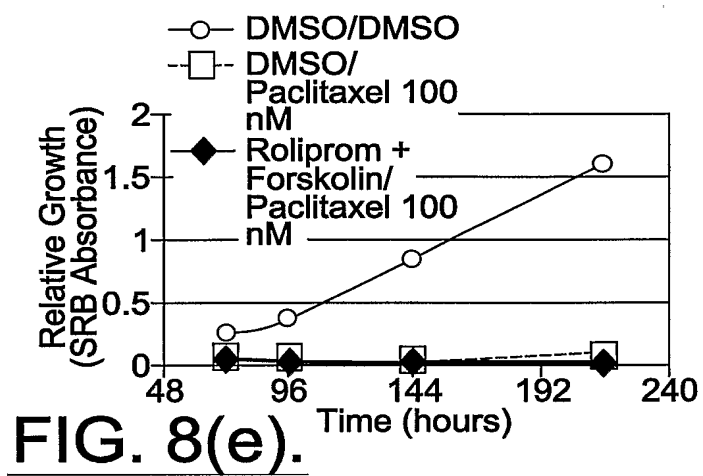
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**FIG. 8a.****FIG. 8(b).**

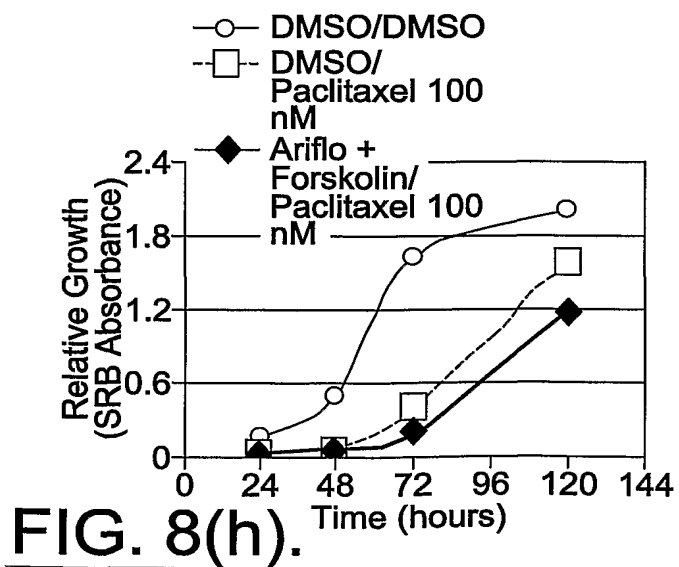
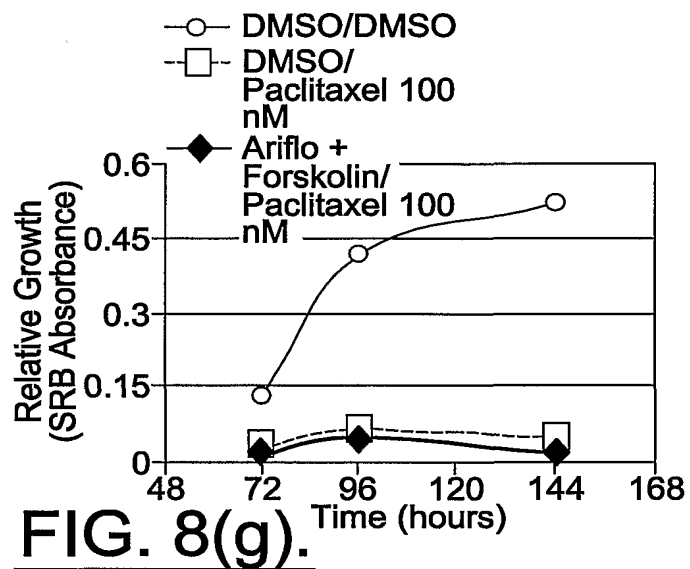
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**FIG. 8(c).****FIG. 8(d).**

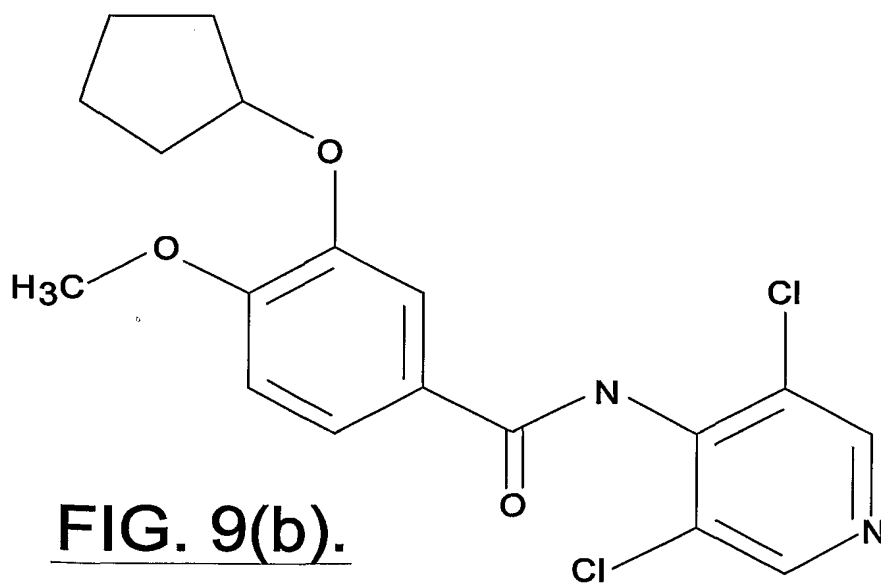
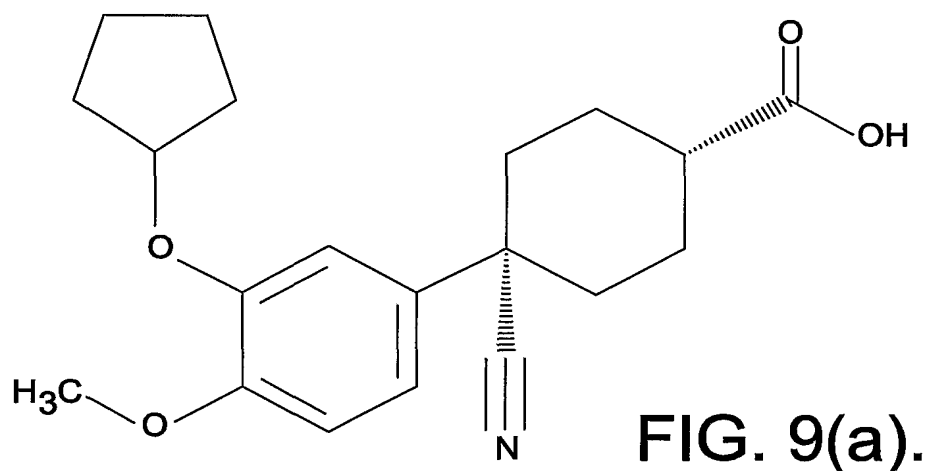
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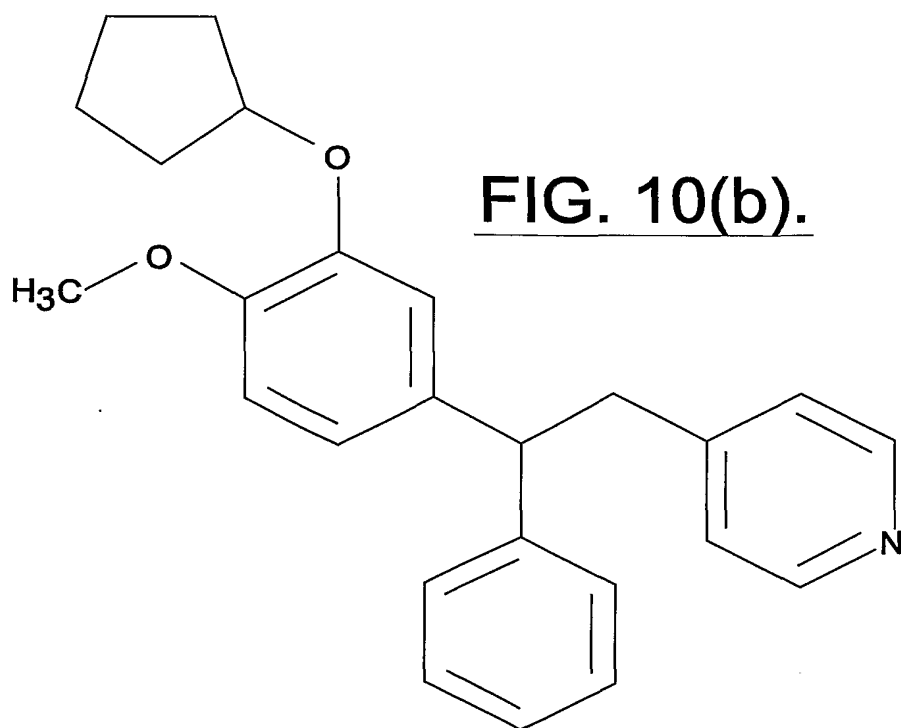
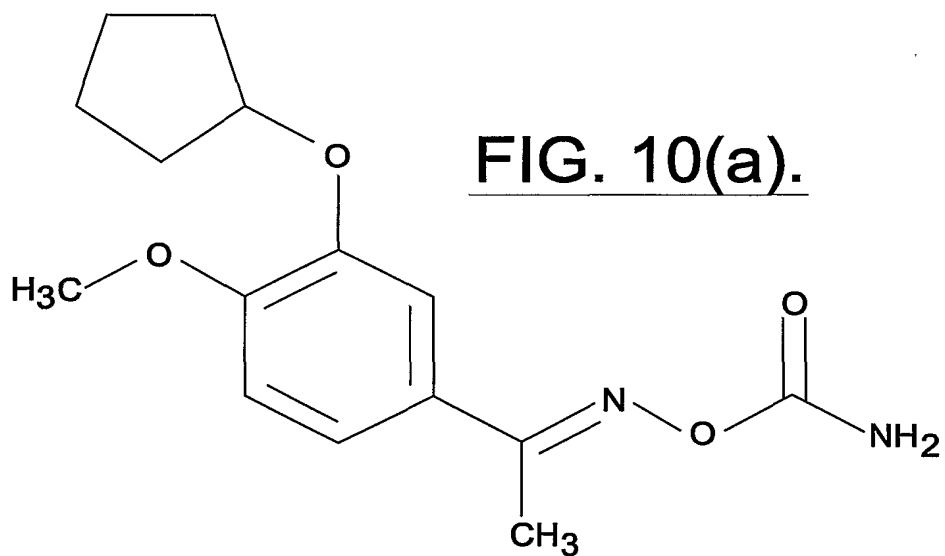
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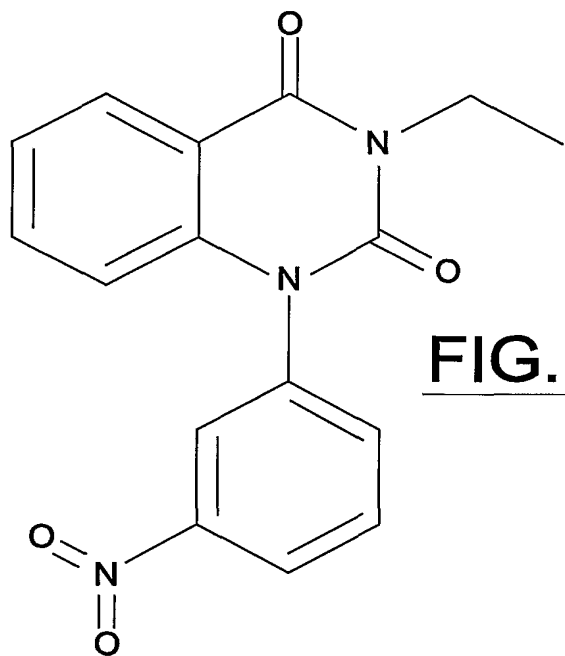
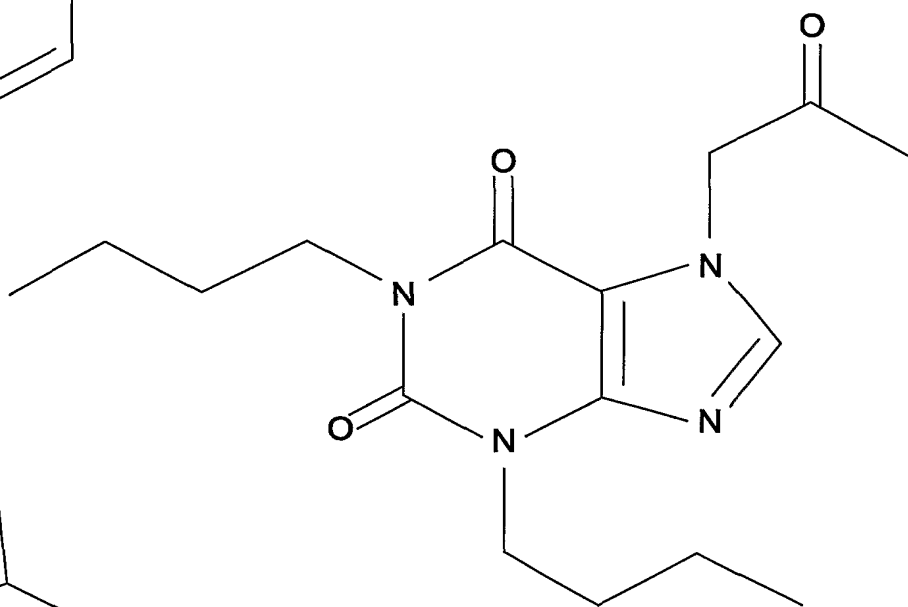
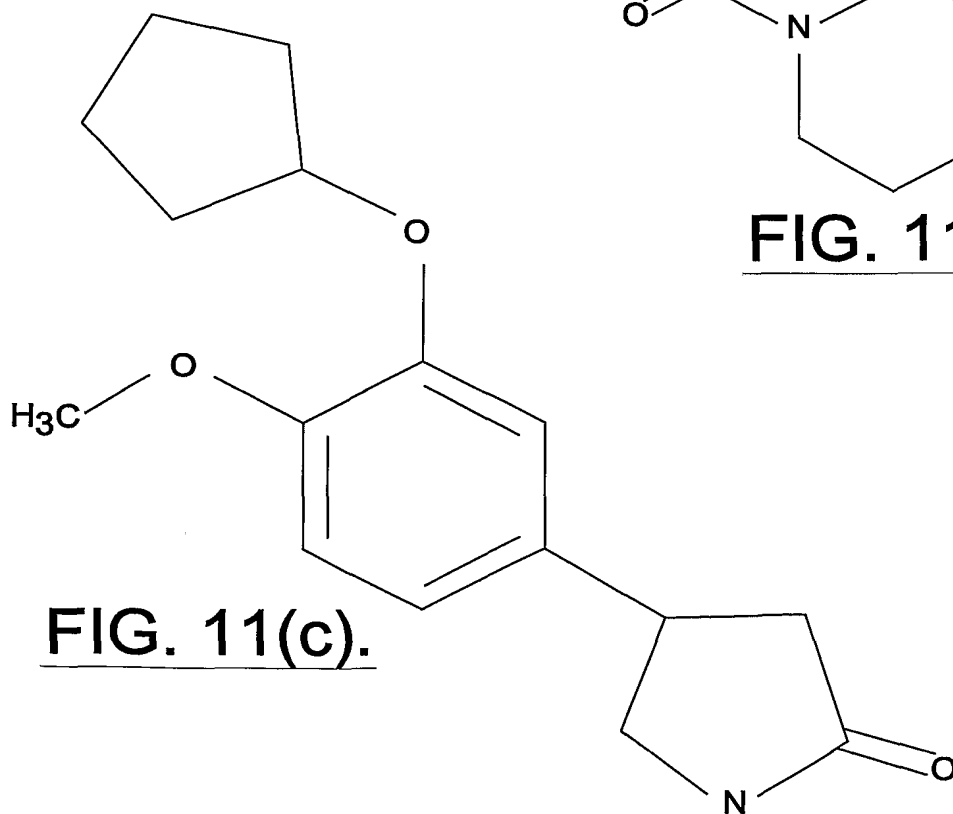
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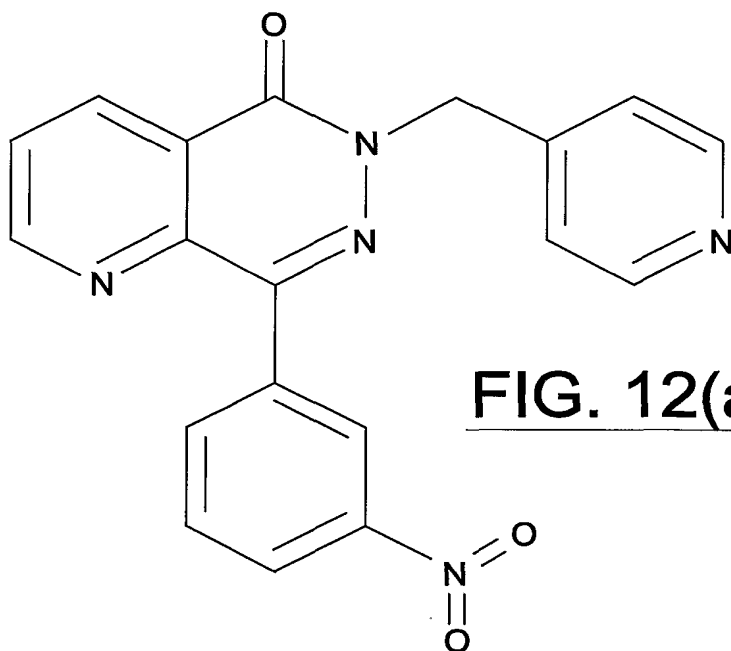
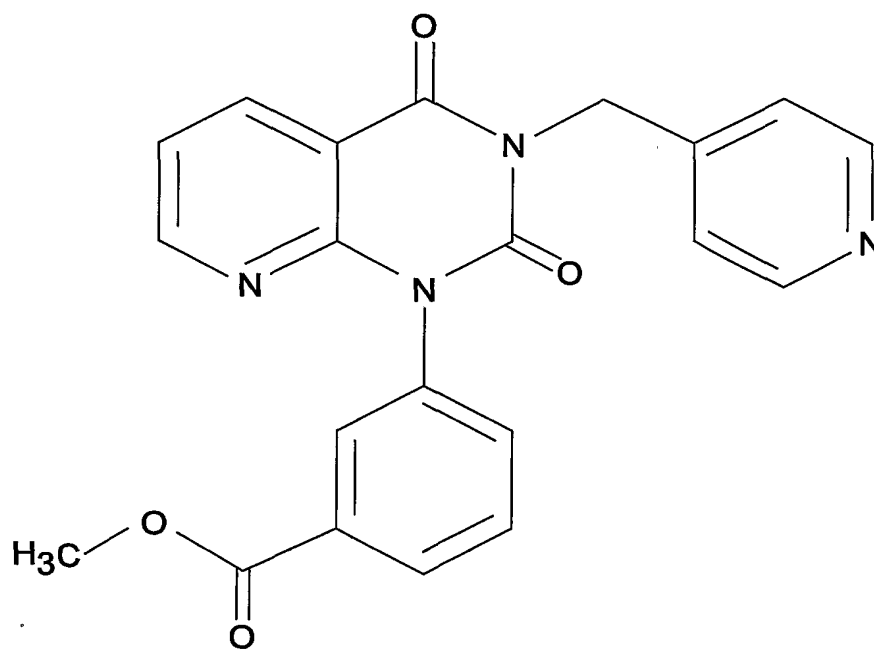
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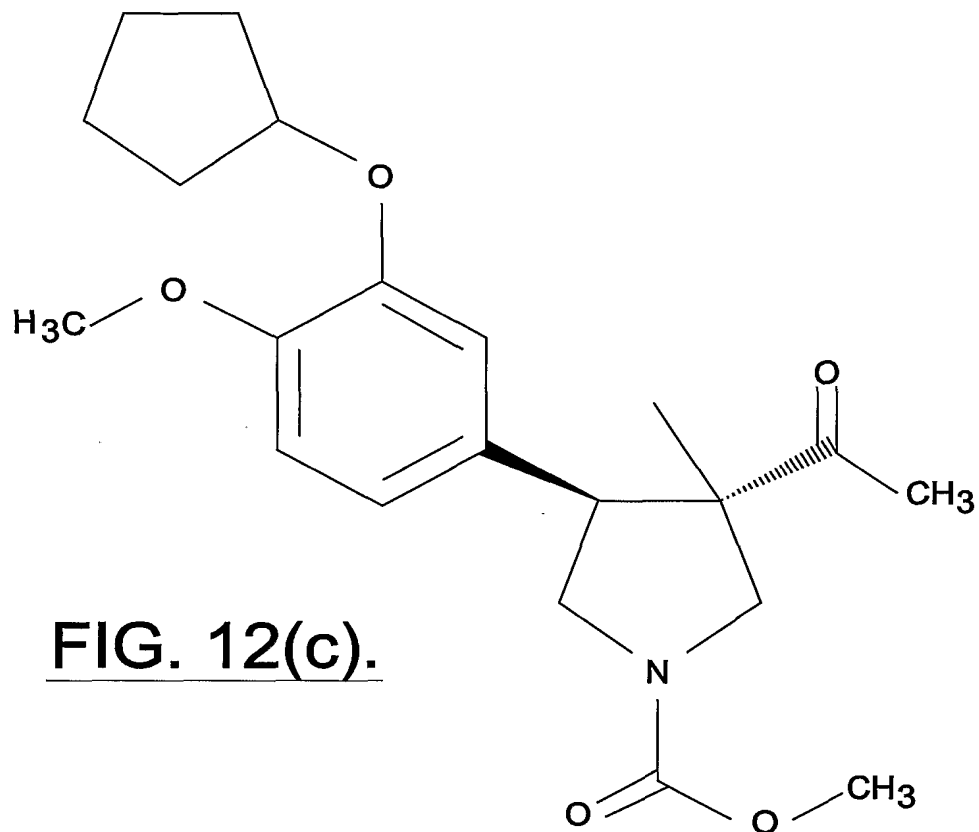
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FIG. 11(a).FIG. 11(b).FIG. 11(c).

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FIG. 12(a).

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FIG. 12(c).

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
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PCT

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A61K 45/06, 31/275, 31/335, 31/47, 31/135, 31/35

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60/209,846 6 June 2000 (06.06.2000) US

(71) Applicant (for all designated States except US): **GLAXO GROUP LIMITED** [GB/GB]; Glaxo Wellcome House, Berkeley Avenue, Greenford, Middlesex UB6 0NN (GB).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **DAVIS, Stephen, Thomas** [US/US]; GlaxoSmithKline, Five Moore Drive, P.O. Box 13398, Research Triangle Park, NC 27709 (US). **DEV, Inderjit, Kumar** [US/US]; GlaxoSmithKline, Five Moore Drive, P.O. Box 13398, Research Triangle Park, NC 27709 (US). **DORNSIFE, Ronna, Ellen** [US/US]; GlaxoSmithKline, Five Moore Drive, P.O. Box 13398, Research Triangle Park, NC 27709 (US). **GRAY, John, Gordon** [US/US]; GlaxoSmithKline, Five Moore Drive, P.O. Box 13398, Research Triangle Park, NC 27709 (US).

(74) Agents: **LEMANOWICZ, John, L.** et al.; GlaxoSmithKline, Five Moore Drive, P.O. Box 13398, Research Triangle Park, NC 27709 (US).

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

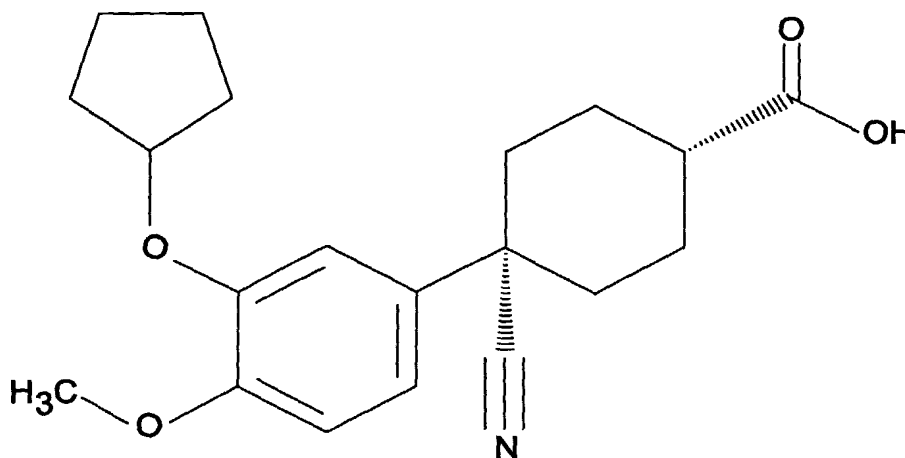
Published:

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

(88) Date of publication of the international search report:
5 December 2002

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: CANCER TREATMENT COMPOSITION CONTAINING AN ANTI-NEOPLASTIC AGENT AND A PDE4 INHIBITOR



(57) Abstract: The invention relates to a method for treating cancer including administering to a mammal therapeutically effective amounts of an anti-neoplastic agent and a PDE4 inhibitor, treatment combinations containing the same as well as a method for attenuating unwanted cancer treatment side effects including myelosuppression and mucositis.



WO 01/093909 A3

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 01/18108

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61P35/00 A61K45/06 A61K31/275 A61K31/335 A61K31/47
A61K31/135 A61K31/35

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

BIOSIS, MEDLINE, CHEM ABS Data, EMBASE, EPO-Internal, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 99 49859 A (MEYSKENS FRANK L JR ;GERNER EUGENE W (US); UNIV ARIZONA (US); UNIV) 7 October 1999 (1999-10-07) page 4, line 4-12; claims 1,6; example 3 ---	1,2,6, 15,20, 29,30, 35-38, 43,48, 49,54
X	WO 00 16621 A (LERNER ADAM ;BOSTON MEDICAL CENTER CORP (US)) 30 March 2000 (2000-03-30) claims 1,7; example 8 --- -/-	1-3,6, 15-17, 20,29, 30,35, 37-40, 43,48, 49,54



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Date of the actual completion of the international search

5 September 2002

Date of mailing of the international search report

02/10/2002

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Gonzalez Ramon, N

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 01/18108

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
E	<p>WO 01 78651 A (CELL PATHWAYS INC) 25 October 2001 (2001-10-25)</p> <p>page 33; figures 6-9; example 2; table 1</p>	<p>1-3,5,6, 15-17, 19,20, 29,30, 35-40, 42,43, 48,49,54</p>
X	<p>EP 1 188 438 A (WARNER LAMBERT CO) 20 March 2002 (2002-03-20) page 3, line 46.47; claims 1,2,6</p>	54
X,P	<p>DORNSIFE RONNA E ET AL: "Phosphodiesterase 4 (PDE4) inhibitors provide selective chemoprotection of human neutrophil progenitors from phase-specific anti-cancer agents with no effect on anti-neoplastic potency." BLOOD, vol. 96, no. 11 Part 1, 16 November 2000 (2000-11-16), page 20a XP001074197 42nd Annual Meeting of the American Society of Hematology; San Francisco, California, USA; December 01-05, 2000 ISSN: 0006-4971 abstract</p>	1-54
X,P	<p>DORNSIFE RONNA E ET AL: "Selective chemoprotection of human neutrophil progenitors and mink lung epithelial cells by phosphodiesterase 4 (PDE4) inhibitors in vitro from the toxicities of phase-specific anti-cancer agents with no effect on anti-neoplastic potency." PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH ANNUAL, vol. 42, March 2001 (2001-03), page 189 XP008007511 92nd Annual Meeting of the American Association for Cancer Research; New Orleans, LA, USA; March 24-28, 2001, March, 2001 ISSN: 0197-016X abstract</p>	1-54

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INTERNATIONAL SEARCH REPORT

International Application No

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
T	PIAZZA G A ET AL: "Exisulind, a novel proapoptotic drug, inhibits rat urinary bladder tumorigenesis." CANCER RESEARCH. UNITED STATES 15 MAY 2001, vol. 61, no. 10, 15 May 2001 (2001-05-15), pages 3961-3968, XP001104534 ISSN: 0008-5472 abstract; figure 4 ----	1-54
A	WO 98 41232 A (BASF AG ;CARTER ADAM (US); SEKUT LES (US); GHAYUR TARIQ (US); TRAC) 24 September 1998 (1998-09-24) page 75, line 13; claims 9,10,12; examples 3,5-8 ----	1-54
A	WO 99 34804 A (UNIV VIRGINIA) 15 July 1999 (1999-07-15) page 12, line 23-26; claims 1,2 -----	1-54

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: partially 1-10,15-24,29,30,33,35-49,52-54

Present claims 1-10,15-24,29,35-48,54 relate to compounds defined by reference to desirable characteristics or properties, namely: "antineoplastic agent which has an adverse effect on normal progenitor cells upon use"; "a PDE4 inhibitor" (claims 1,15,29,38,48,54); "phase specific anti-neoplastic agent" (claim 2,16,39); "agent effective in raising intracellular concentrations of cAMP or analogs" (claim 9,23,33,46,52); The claims cover all compounds having these characteristics or properties, whereas the application provides support within the meaning of Article 6 PCT and disclosure within the meaning of Article 5 PCT for only a very limited number of such compounds. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible.

Independent of the above reasoning, the claims also lack clarity (Article 6 PCT). An attempt is made to define the compounds by reference to a result to be achieved. Again, this lack of clarity in the present case is such as to render a meaningful search over the whole of the claimed scope impossible.

Moreover claims 3, 6, 17, 20,30,40, 43,49 relate to an extremely large number of possible compounds defined as "diterpenoids", "vinca alkaloids", "epipodo-phyllotoxins", "antimetabolites", "camptothecins" (claims 3,17,40) or as a Markush structures of claims 6,20,30,43,49. In fact, said claims contain so many options, variables, possible permutations that a lack of clarity (and conciseness) within the meaning of Article 6 PCT arises to such an extent as to render a meaningful search of the claims not fully possible.

Consequently, the search has been carried out for those parts of the claims which appear to be clear, supported and disclosed, namely those parts relating to the compounds/compositions of claims 11-14,25-28,31-32,34,50,51 and those mentioned in the examples and the combination of antineoplastic agents specifically worded in claim 4,5,18,19,41,42 (namely paclitaxel and topotecan) with the PDE4 inhibitors listed in claims 7,8,21,22,31,32,44,45 and optionally the agent of claims 10,24,47,53.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 01/18108

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Although claims 1-14,29-37,54 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☒ Claims Nos.: partially 1-10,15-24,29,30,33,35-49,52-54
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 01/18108

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